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**PDVE**

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**UKKI**

OY 61 KGLEPAAIFHLKLPAINDTDHGCLIGFNETSCKLKLDGFEFEVLFKFTTEFGKSVI 120  
 DB 121 NVDVVELLTKLGMIDQELNKLTKTHSPKFDRLGLGRLOGLKTYWRHFASTVLSAM 180  
 OY 121 NVDVVELLTKLGMIDQELNKLTKTHSPKFDRLGLGRLOGLKTYWRHFASTVLSAM 180  
 DB 181 EKFAQAVRVLDSDPDVTPDVHDK 204  
 OY 181 EKFAQAVRVLDSDPDVTPDVHDK 204  
 RESULT 2  
 ID W40103 standard; Protein; 204 AA.  
 AC W40103;  
 DT 15-JUL-1998 (first entry)  
 DE Human herpesvirus 8 (HHV-8) interleukin-6.  
 KW Bcl-2 homologue; IE-1A; IE-1B; viral macrophage inhibitory protein;  
 KM Kaposi's sarcoma; delta-chemokine-like; amplification; PCR; VIL-6.  
 OS WO9804284-A1.  
 PN 05-FEB-1998.  
 PR 24-JUL-1997: U12931.  
 PA (U120 ) UNIT JOHNS HOPKINS.  
 PI Hardwick JM, Hayward GS, Nicholas J, Reitz MR;  
 PT WPI; 98-130422/12.  
 PT New human herpes virus gene region containing 8 open reading frames  
 PT - useful for, e.g. diagnosing Kaposi's sarcoma or body cavity based  
 PT large cell lymphoma  
 PS Claim 1: Pages 55-60; 84pp; English.  
 CC The sequence represents a novel human herpesvirus 8 (HHV-8)  
 CC interleukin-6. The invention claims for novel genes, which includes  
 CC the viral interleukin-6 gene, found at the divergent DL-B locus. HHV-8  
 CC divergent locus DL-B lies between open reading frames 11 and 17.  
 CC Sequencing of the HHV-8 divergent locus DL-B revealed the presence of  
 CC nine viral ORFs with gene products related to cellular proteins. These  
 CC proteins include the thymidylate synthase (TS, W40100), dihydrofolate  
 CC reductase (DHFR, see W40101), Bcl-2 homologue (W40102), IE-1A (W40107),  
 CC IE-1B (W40108) and, four cytokines which include viral interleukin-6  
 CC (vIL-6), viral macrophage inhibitory protein (vMIP)-1A (W40104) and -1B  
 CC (W40105) and beta-chemokine-like (BCK, W40106) protein. The invention  
 CC claims the mentioned proteins and a polynucleotide containing HHV-8  
 CC genes encoding one or more of these proteins. The invention also claims  
 CC that the polynucleotide and the proteins may be used directly or  
 CC indirectly, e.g. using antibodies to the proteins, to diagnose an HHV-8  
 CC associated disease, e.g. Kaposi's sarcoma, Castleman's disease, multiple  
 CC myeloma and body cavity based large cell lymphoma (BCBL). The proteins  
 CC have also been claimed to be useful in screening compounds for drugs to  
 CC treat HHV-8 diseases.  
 SQ Sequence 204 AA;

Query Match 100.0%; Score 1579; DB 1; Length 204;  
 Best Local Similarity 100.0%; Pred. No. 9, 31e-140;  
 Matches 204; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 1 MCFKRLMSLLVGSILVSGTRGKLPDAPFEKDLIQRNLNMLWVIDECFRLCYRTGIC 60  
 OY 1 MCFKRLMSLLVGSILVSGTRGKLPDAPFEKDLIQRNLNMLWVIDECFRLCYRTGIC 60  
 DB 61 KGLEPAAIFHLKLPAINDTDHGCLIGFNETSCKLKLDGFEFEVLFKFTTEFGKSVI 120  
 OY 61 KGLEPAAIFHLKLPAINDTDHGCLIGFNETSCKLKLDGFEFEVLFKFTTEFGKSVI 120  
 DB 121 NVDVVELLTKLGMIDQELNKLTKTHSPKFDRLGLGRLOGLKTYWRHFASTVLSAM 180  
 OY 121 NVDVVELLTKLGMIDQELNKLTKTHSPKFDRLGLGRLOGLKTYWRHFASTVLSAM 180  
 DB 181 EKFAQAVRVLDSDPDVTPDVHDK 204  
 OY 181 EKFAQAVRVLDSDPDVTPDVHDK 204

RESULT 3  
 ID W23944 standard; Protein; 204 AA.  
 AC W23944;  
 DT 30-JUN-1998 (first entry)  
 DE Human herpesvirus 8 interleukin-6.  
 KW Interleukin-6; IL-6; human herpesvirus 8; HHV8; shotgun-cloning;  
 KM Antibody; diagnosis; treatment.  
 OS Human herpesvirus 8.  
 PN WO9803657-A1.  
 PR 29-JAN-1998.  
 PR 19-JUL-1996: E03199.  
 PA (BEHW ) BEHRING DIAGNOSTICS GMBH.  
 PI Albrecht J, Fleckenstein B, Friedman-Klen A, Huang Y,  
 PI Neipel F;  
 PT WPI; 98-120781/11.  
 PT Viral interleukin-6 produced recombinantly from human herpes virus 8  
 PT DNA - and related nucleic acid and antibodies, used for diagnosis  
 PT and treatment of herpes 8 infection or related diseases, e.g. Kaposi  
 PT sarcoma  
 PS Claim 2: Fig 2: 19pp; English.  
 CC The interleukin-6 (IL-6) and protein W23944 derived from the sequence  
 CC can be used to detect antibodies and antibodies can be used to detect  
 CC IL-6. This can be used for the diagnosis of HHV8 infection or  
 CC associated diseases such as Kaposi sarcoma or kidney cell carcinoma.  
 CC Antibodies, proteins and the gene sequence can all be used in the  
 CC treatment of infections and diseases as mentioned above.  
 SQ Sequence 204 AA;

Query Match 100.0%; Score 1579; DB 1; Length 204;  
 Best Local Similarity 100.0%; Pred. No. 9, 31e-140;  
 Matches 204; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 1 MCFKRLMSLLVGSILVSGTRGKLPDAPFEKDLIQRNLNMLWVIDECFRLCYRTGIC 60  
 OY 1 MCFKRLMSLLVGSILVSGTRGKLPDAPFEKDLIQRNLNMLWVIDECFRLCYRTGIC 60  
 DB 61 KGLEPAAIFHLKLPAINDTDHGCLIGFNETSCKLKLDGFEFEVLFKFTTEFGKSVI 120  
 OY 61 KGLEPAAIFHLKLPAINDTDHGCLIGFNETSCKLKLDGFEFEVLFKFTTEFGKSVI 120  
 DB 121 NVDVVELLTKLGMIDQELNKLTKTHSPKFDRLGLGRLOGLKTYWRHFASTVLSAM 180  
 OY 121 NVDVVELLTKLGMIDQELNKLTKTHSPKFDRLGLGRLOGLKTYWRHFASTVLSAM 180  
 DB 181 EKFAQAVRVLDSDPDVTPDVHDK 204  
 OY 181 EKFAQAVRVLDSDPDVTPDVHDK 204

RESULT 4  
 ID W95015 standard; peptide; 185 AA.  
 AC W95015;  
 DT 21-MAY-1999 (first entry)  
 DE Kaposi's sarcoma herpes virus interleukin-6 (IL-6) polypeptide.  
 KW Cytokine; interleukin-B30; IL-B30; forensic science; cell proliferation;  
 OS Kaposi's sarcoma herpes virus.  
 PN WO9905280-A1.  
 PR 04-FEB-1999.  
 PR 24-JUL-1998: U15423.  
 PR 25-JUL-1997: US-900905.  
 PA (SCHE ) SCHERING CORP.  
 PI Bazan JF;  
 PT WPI; 99-142935/12.  
 PT Newly isolated or recombinant polynucleotide encoding mammalian  
 PT cytokine interleukin-B30 (IL-B30), including fragments - useful for  
 PT regulating activation, development, differentiation and function of  
 PT various cell types, and for diagnosing and treating conditions  
 PT associated with IL-B30  
 PS Disclosure; Page 11-12; 83pp; English.

PT	factor to identify binding
PS	Claim 8; : 26pp; English.

sequence 184 AA;

Query Match	16.7%;	Score 264;	DB 1;	Length 184;
Best Local Similarity	27.7%;	Pred. No. 2.47e-13;		
Matches	39.	Conservative	39.	Mismatches 63.
				Indels 0.
				Gaps 0.









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FH Key Location/Qualifiers
FT Peptide 1..28
FT Protein /note="signal peptide"
FT Protein 29..212
FT Misc_difference 82 /note="mature protein"
FT FT /label=K82P
FT FT /note="wild type lys is replaced by pro"
FT Misc_difference 187 /label=Q187E
FT FT /note="wild type Gln is replaced by Glu"
FT Misc_difference 190 /label=T190P
FT FT /note="wild type Thr is replaced by Pro"
FT Misc_difference 198 /label=F198L
FT FT /note="wild type Phe is replaced by Leu"
FT Misc_difference 204 /label=S204R
FT FT /note="wild type Ser is replaced by Arg"
FT FT
FT W09738103-A1.
PD 16-OCT-1987.
PF 09-APR-1996; E01506.
PR 09-APR-1996; W0-E01506.
PA (ISTF ) ARS APPLIED RES SYSTEMS HOLDING NV.
PI Ehlers M, Grotzinger J, Rose-John S;
DR WPT: 97-512720/47.
PT New interleukin-6 mutein polypeptides - useful as IL-6 antagonists
PT for treating e.g. plasmacytoma/myeloma, osteoporosis and neoplastic
PT and autoimmune diseases.
PS Claim 1: Pages 15-16; 38pp; English.
CC This is a interleukin-6 (IL-6) mutein polypeptide. This IL-6 mutein
CC comprises a mature protein of 184 amino acids. This polypeptide is
CC created by point mutations on the wild-type human IL-6, at positions
CC 34, 159, 162, 170, 176 of the mature human IL-6. The DNA sequence encoding
CC this IL-6 mutein and the sequences encoding variants having the same
CC activity resulting from the degeneracy of the genetic code or point
CC mutations can be used to transform a host cell. The IL-6 mutein can act
CC as IL-6 antagonist. This polypeptide and its fragments can be used for
CC treating diseases in which IL-6 has a pathogenic action such as
CC plasmacytoma myeloma, osteoporosis and neoplastic and autoimmune
CC diseases.
SQ Sequence 212 AA;

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Query Match 16.2% Score 256; DB 1; Length 212;
Best Local Similarity 27.7% Pred.No.1.28e-12;
Matches 39; Conservative 36; Mismatches 64; Indels 0; Gaps 0;

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DB 69 KETCNKSNMCESSPALAENNINLPKMAEKDCFOGFEENETCLVKITGLLEFEVLEY 128
OY 51 RDLGRTGICIGILEPAAIFHLKLPAINDTDHGCLIGFNETSCLKRLADGFEFEVLEPF 110
DB 129 LONRESESEQARAVQMSKVIQFLQKKAKNLDATTPDPTNMSLTKLQANQWLED 188
OY 111 LTTREGKSYINADVWMLLTGTLGMDIQEBLNKLTYSPPKFDRLGLRLQGLKLVWRH 170
DB 189 MPTHLIRSLKEFLQSLRAL 209
OY 171 PASFYLSAMEKFAQAGAVRVL 191

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Search completed: Fri Sep 15 16:32:05 2000
Job time : 12 secs.

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[illegible]

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##residues      28-40 ##label MAY3
##experimental_source FS-4 fibroblasts
##note          sequence extracted from NCBI backbone (NCBIP:63787)
##note          this 23-25k form contained O-linked but not N-linked
##note          carbohydrate

REFERENCE
#authors        JX0305
#journal        Orita, T.; Ohneda, M.; Hasegawa, M.; Kubonikwa, H.; Esaki, K.:
#title          Ochi, N.
#cross-references J. Biochem. (1994) 115:345-350
#contents       Polypeptide and carbohydrate structure of recombinant human
                  interleukin-6 produced in chinese hamster ovary cells.
                  MIMD:94266765
#contents       annotation; modified sites in recombinant protein from CHO
                  cells
REFERENCE
S04981

Note: remainder of annotations omitted.

Query Match      16.0%: Score 252; DB 1; Length 212;
Best Local Similarity 27.7%: Pred. No. 2,82e-25;
Matches 39; Conservative 37; Mismatches 65; Indels 0; Gaps 0;

Db 69 KETCNMNCSSSEKALAEENLNLPKAAEKDCFCOSFNEETCKVITITGLTEFEVLEYL 128
QY 51 RDLCTRGICIGILEPAIPIHKLPAINDRDGCLGFCFNETSCLKLADGFFFEVLFKF 110
Db 129 LQNFESSEQARAQVQNSTKVLQIQLQKAKNDATITPPPTNASLITRLQANQWQLQD 188
QY 111 LTTEFGKSVINQVDELTKLTGLMDIOELNKLTKTHYSPKFRDGLGLQGLKQVWRH 170
Db 189 MTHLILRSFKFELQSSLRAL 209
QY 171 FASFYLSAMEKFAQGAQVRL 191

RESULT 4
ENTRY      T09216 #type complete
TITLE      interleukin-6 precursor - horse
ORGANISM   #format_name Equus caballus #common_name domestic horse
DATE       11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change
          23-Jul-1999

ACCESSIONS T09216
REFERENCE   Z16613
#authors    Swiderski, C.E.; Horohov, D.W.
#submission submitted to the EMBL Data Library, July 1996
#accession  T09216
#status     preliminary; translated from GB/EMBL/DBJ
#molecule_type mRNA
#residues   1-208 ##label SWI
#cross-references EMBL:064794; NID:g2654387; PID:g2654388

GENETICS
#gene        IL-6
CLASSIFICATION
KEYWORDS     #superfamily interleukin-6
              cytokine; growth factor
SUMMARY      #length 208 #molecular_weight 23419 #checksum 3370

Query Match      15.1%: Score 238; DB 2; Length 208;
Best Local Similarity 27.3%: Pred. No. 9.30e-23;
Matches 36; Conservative 37; Mismatches 58; Indels 1; Gaps 1;

Db 67 EMCNNSKSCNSKEVLAENNLNLPKAAEKDCFCOSGFNETCLMKITITGSEFOIYLEYL 126
QY 52 DCLYRTGICIGILEPAIPIHKLPAINDTDHCGILIGNENISCLKLADGFFFEVLEKFL 111
Db 127 QNEFGKEKENTKTOISKVLV-QILQKAKNPVYITPPDTAKSSSLAKHSQNEHAKNT 185
QY 112 TTEFGKSVINQVDELTKLTGLMDIOELNKLTKTHYSPKFRDGLGLRQGLKQVWRH 171
Db 186 TTHLILRSLEDF 197
QY 172 ASFYLSAMEKF 183

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RESULT      5
ENTRY       S29549      #type complete
TITLE      Interleukin-6 - sheep
ORGANISM   #formal_name Ovis orientalis aries, Ovis ammon aries
           #common_name domestic sheep
DATE       10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change
           10-Sep-1999
ACCESSIONS S29549
REFERENCE   S29549
#authors   Ebrahim, B.
#submission Submitted to the EMBL Data Library, October 1992
#accession S29549
#status    preliminary
#molecule_type mRNA
#residues  1-208 ##label EBR
#cross-references EMBL:X68723
CLASSIFICATION
#superfamily Interleukin-6
SUMMARY    #length 208 #molecular_weight 23526 #checksum 7927

Query Match      14.1% Score 222; DB 1; Length 208;
Best Local Similarity 27.1%; Pred. No. 6,428-20;
Matches 36; Conservative 37; Mismatches 56; Indels 4; Gaps 4;

Db 69 KEICRNDCEKSKETLAENKLTLPKMEKQCGQSGFNAVCLIKTTAGLLEQYIDF 128
   :::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Oy 51 RDLCYRTGICGILEPAIIFHLKLPAINDTDHGILGFNETSCKLADGFEFEVLEKF 110
   :::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|

Db 129 LONFE-GNQT-VWELQ-SI-RTLIQLEKINAGLITTPATHRDLEKQSSNEWYKN 184
   :::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Oy 111 LTTEGKSVINVDVETLTLTGMDIOBELNKLTKRTHSPRFNGLGRIGKGYVVRH 170
   :::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|

Db 185 AKVITLRLENF 197
Oy 171 FASFYVLSAMERF 183

RESULT      6
ENTRY       ICMS6      #type complete
TITLE      Interleukin-6 precursor - mouse
ALTERNATE_NAMES B-cell hybridoma growth factor; B-cell stimulating factor 2;
                hepatocyte-stimulating factor; IL-6; Interferon beta-11;
                Interleukin-HP1; myeloid differentiation inducer MGI-2A;
                plasmacytoma growth factor
ORGANISM   #formal_name Mus musculus #common_name house mouse
DATE       30-Jun-1990 #sequence_revision 30-Jun-1990 #text_change
           27-Jun-1999
ACCESSIONS A30531; A27610; A30571; S01323; S12103; E34047; A26662;
           A40486; A60799; S10241; S38254
REFERENCE   A30531
#authors   Tanabe, O.; Akira, S.; Kamitaya, T.; Wong, G.C.; Hirano, T.;
           Kishimoto, T.
#journal   J. Immunol. (1988) 141:3875-3881
#title     Genomic structure of the murine IL-6 gene. High degree
           conservation of potential regulatory sequences between
           mouse and human.
#cross-references MUID:89035525
#accession A30531
#molecule_type DNA
#residues  1-211 ##label TAN
#cross-references GB:M20572; NID:G198369; PIDN:AAA39302.1; PID:9387386
REFERENCE   A27610
#authors   Van Snick, J.; Cayphas, S.; Szikora, J.P.; Renauld, J.C.; Van
           Roost, E.; Boon, T.; Simpson, R.J.
#journal   Eur. J. Immunol. (1988) 18:193-197
#title     CDNA cloning of murine Interleukin-HP1: homology with human
           Interleukin 6.
#cross-references MUID:88166883
#accession A27610
#molecule_type mRNA
#residues  1-211 ##label VAN
#cross-references GB:X06203; NID:952701; PIDN:CAA29560.1; PID:952702
REFERENCE   A30571
#authors   Mock, B.A.; Nordan, R.P.; Justice, M.J.; Kozak, C.; Jenkins,

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#journal    N.A.; Copeland, N.G.; Clark, S.C.; Wong, G.C.; Rudikoff, S.
#title      J. Immunol. (1989) 142:1372-1376
#cross-references MUID:89124383
#accession  A30571
#molecule_type mRNA
#residues  5-211 ##label MOC
#cross-references GB:M44221; NID:9341131; PIDN:AAA68814.1; PID:9870699
REFERENCE   S01323
#authors   Simpson, R.J.; Moritz, R.L.; Rubira, M.R.; Van Snick, J.
#journal   Eur. J. Biochem. (1988) 176:187-197
#title     Murine hybridoma/plasmacytoma growth factor. Complete
           amino-acid sequence and relation to human Interleukin-6.
#cross-references MUID:88329055
#accession S01323
#molecule_type protein
#residues  25-166,'X',168-211 ##label SIM
#note       The sequence from Fig. 11 is inconsistent with that from
           Fig. 10 in having 103-Asn

REFERENCE   S12103
#authors   Grenett, H.E.; Fuentes, N.L.; Fuller, G.M.
#journal   Nucleic Acids Res. (1990) 18:6455
#title     Cloning and sequence analysis of the cDNA for murine
           Interleukin-6.
#cross-references MUID:91057159
#accession S12103
#molecule_type mRNA
#residues  1-211 ##label GRE
#cross-references EMBL:X54542; NID:952727; PIDN:CAA38411.1; PID:952728
REFERENCE   A90157
#authors   Jahnen, W.; Ward, L.D.; Reid, G.E.; Moritz, R.L.; Simpson,
           R.J.
#journal   Biochem. Biophys. Res. Commun. (1990) 166:139-145
#title     Internal amino acid sequencing of proteins by in situ
           cyanogen bromide cleavage in polyacrylamide gels.
#cross-references MUID:90147691
#accession E34047
#molecule_type protein
#residues  56-65,'X',71-75;78-94;128-148 ##label JAS
REFERENCE   A26662
#authors   Van Snick, J.; Cayphas, S.; Yank, A.; Uyttenhove, C.; Couille,
           P.G.; Rubira, M.R.; Simpson, R.J.
#journal   Proc. Natl. Acad. Sci. U.S.A. (1988) 83:9679-9683
#title     Purification and NH2-terminal amino acid sequence of a
           T-cell-derived lymphokine with growth factor activity for
           B-cell hybridomas.
#cross-references MUID:87092311
#accession A26662
#molecule_type protein
#residues  25-39,'X',41-42,'X',44-45 ##label VSN
REFERENCE   A40486
#authors   Chu, C.P.; Moulds, C.; Coffman, R.L.; Rennick, D.; Lee, F.
           Proc. Natl. Acad. Sci. U.S.A. (1988) 85:7099-7103
#title     Multiple biological activities are expressed by a mouse
           Interleukin 6 cDNA clone isolated from bone marrow stromal
           cells.
#cross-references MUID:89017145
#accession A40486
#molecule_type mRNA
#residues  1-211 ##label CHI
#cross-references GB:J03783; NID:G198367; PIDN:AAA39301.1; PID:9309410
REFERENCE   A60799
#authors   Shabo, Y.; Lotem, J.; Rubinstein, M.; Revel, M.; Clark, S.C.;
           Wolf, S.F.; Kamen, R.; Sachs, L.
#journal   Blood (1988) 72:2070-2073
#title     The myeloid blood cell differentiation-inducing protein
           MGI-2A is interleukin-6.
#cross-references MUID:89062753
#accession A60799
#molecule_type protein
#residues  77-98 ##label SHA
REFERENCE   S10241

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#authors          J. Blankenstein, T.; Qin, Z.; Li, W.; Diamantstein, T.
#journal          J. Exp. Med. (1990) 171:965-970
#title            DNA rearrangement and constitutive expression of the
                  interleukin 6 gene in a mouse plasmacytoma.
#cross-references MIMD:90171860
#accession        S10241
#status            preliminary
##molecule_type  DNA
##residues         1-6 ##label BJA
##cross-references EMBL:X51457; NID:g49738; PIDN:CAA35824.1; PID:g581860
REFERENCE
#authors          Zhang, J.G.; Reid, G.E.; Moritz, R.L.; Ward, L.D.; Simpson,
                  R.J.
#journal          Eur. J. Biochem. (1993) 217:53-59
#title            Specific covalent modification of the tryptophan residues in
                  murine interleukin-6. Effect on biological activity and
                  conformational stability.
#cross-references MIMD:9403075
#accession        S38254
#status            preliminary
##molecule_type  protein
##residues         38-60;75,'X','77-79;176-203 ##label ZHA
GENETICS
#gene              IL-6
#map_position      7/1: 68/3; 106/3; 156/3
#introns           1
CLASSIFICATION
KEYWORDS           *superfamily interleukin-6
                  Castleman's disease; cytokine; growth factor;
                  immunoregulation; lymphokine; macrophage; rheumatoid
                  arthritis
FEATURE
1-24               #domain signal sequence #status predicted #label SIG\
SUMMARY            25-211 #product interleukin-6 #status experimental #label MAT
                  #length 211 #molecular-weight 24384 #checksum 5652
Query Match       14 13; Score 223; DB 1; Length 211;
Best Local Similarity 26.7%; Pident. No. 4; 28e-20;
Matches 40; Conservative 42; Mismatches 65; Indels 3; Gaps 3;
Db 58 VLMEIVEMREKLENGNSDCMNNDALAEENNLKLPETQRNDGCGYGTGYNOECLIKRISGL 117
   :|||:::||: ||: ||: ||||| | | | | | | | | | | | | | | | | | | | |
Qy 42 MLMWIDCEPFDLCRGICIGILEPAIRIHLKIPAINOTDHGGLIGFNETSKKLACGF 101
   :::::|||||::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Db 118 LEVHSYLEYMKNNLKDKKKARVLRDTFTLLIHFNQEVKKLRIIVLPTP-ISNALITD 176
   :|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Qy 102 FEEFEVLKFELTFEF-GKSYINVDVMELLTKTGMDIOELRLKTRTHSPRFDRGLT-G 159
   :|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Db 177 KLSQKEMLRKTRTIQTPIKSLSEFLAVTLR 206
   :|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Qy 160 RLQGLKIWRHFPASFYLAMEKFAQQAVR 189
RESULT
ENTRY             A56610 #type complete
TITLE             Interleukin-6 precursor - bovine
ORGANISM          Homo sapiens
DATE              10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change
ACCSSIONS         A56610: s22162
REFERENCE         A56610
AUTHORS           Drocognans, L.; Cludts, I.; Cleuter, Y.; Kettmann, R.; Burny,
                  A.
#journal          DNA Seq. (1992) 2:411-413
#title            Nucleotide sequence of bovine interleukin-6 cDNA.
#cross-references MIMD:93076003
#accession        A56610
#status            preliminary
##molecule_type  mRNA
##residues         1-208 ##label DR0
##cross-references EMBL:X57317; NID:g2193; PIDN:CAA40572.1; PID:g2194
#experimental_source BLV induced B cell-Lymphosarcoma
#note             *sequence extracted from NCBI Backbone (NCBIF:118917)
CLASSIFICATION    #superfamily interleukin-6

```

KEYWORDS	cytokine		
SUMMARY	#length 208	#molecular-weight 23758	#checksum 8010
Query Match	12.8%;	Score 202;	DB 1; Length 208;
Best Local Similarity	23.2%;	Pred. No. 1,93e-16;	
Matches	33;	Conservative	48; Mismatches 55; Indels 6; Gaps 6;
Db	69	KEICEKNDDESSSEKTELAENKLNLPMEKEDGCFOSGFNOAICLRITAGILEYQIYIDY	128
QY	51	RDLCYRFGICIGLIEPAIAFHLEKLPAINDTDHGCLIGFNETSCLEKKLADGFEFEVLFKE	110
Db	129	LQNEYE-GNCE-VNRDL-RKNI-RTLIQTLKKIKINDLITPAT-NTDLEKMOSSNEVWK	183
QY	111	LTTEEGKSVIYNVMELETTFTLQMDIOEELN-KLTKTHYSPEKFDGILGRIOGLKTIWVR	169
Db	184	NAKIIILIRNLLENFLOFSLRAI	205
QY	170	HFAFYILSAMKFAAGAVRVL	191
RESULT	8		
ENTRY	A34247	#type complete	
TITLE	interleukin-6 precursor - rat		
ALTERNATE_NAMES	IL-6		
ORGANISM	#formal_name Rattus norvegicus #common_name Norway rat		
DATE	15-Jun-1990 #sequence_revision 15-Jun-1990 #text_change 16-Jul-1999		
ACCESSIONS	A34247		
REFERENCE	A34247		
#authors	Northemann, W.; Brackik, T.A.; Hattori, M.; Lee, F.; Fey, G.H.		
#journal	J. Biol. Chem. (1989) 264:16072-16082		
#title	Structure of the rat interleukin 6 gene and its expression in macrophage-derived cells.		
#cross-references	MIMD:89380206		
#accession	A34247		
#status	preliminary		
#molecule_type	mRNA		
#residues	1-211 #label NOR		
#cross-references	GB:M6744; NID:9204915; PIDN:AAAT7659.1; PID:9204916		
CLASSIFICATION	#superfamily Interleukin-6		
KEYWORDS	cytokine; growth factor; immunoregulation; lymphokine; macrophage		
SUMMARY	#length 211	#molecular-weight 24357	#checksum 5864
Query Match	12.0%;	Score 189;	DB 2; Length 211;
Best Local Similarity	25.7%;	Pred. No. 3.15e-14;	
Matches	39;	Conservative	40; Mismatches 72; Indels 1; Gaps 1;
Db	55	ITTVILREILREKRELCNGNSDCNNSDALSENNKLPEIQNRDGCFGYGNQELCLKIC	114
QY	39	LNMLAWIDECFRDLCYRGICIGLIEPAIAFHLEKLPAINDTDHGCLIGFNETSCLKTLA	98
Db	115	SGLEEFYFLFYKANNLQDNKKDKANVIOSTNTELVHIFKQEIKNYSYIVLPTPSNALL	174
QY	99	DGFEFEVLFKFLFTEEGKSVIYNVD-VMELEKTRFTLQMDIOEELNKLTKTHYSPEKFDGIL	157
Db	175	MEKLESOKEMLRRTKTIQLILKALEFLKXTMR	206
QY	158	LGRIOGLKTIWVRHFAFYILSAMKFAAGAVR	189
RESULT	9		
ENTRY	I46084	#type complete	
TITLE	interleukin 6 cat		
ORGANISM	#formal_name Felis silvestris catus #common_name domestic cat		
DATE	16-Aug-1996 #sequence_revision 16-Aug-1996 #text_change 16-Jul-1999		
ACCESSIONS	I46084		
REFERENCE	I46084		
#authors	Bradley, W.G.; Gibbs, C.; Kraus, L.; Good, R.A.; Day, N.K.		
#journal	Proc. Soc. Exp. Biol. Med. (1993) 204:301-305		
#title	Molecular cloning and characterization of a cDNA encoding		

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#feline.interleukin-6.
#cross-references MUID:94052249
#accession I46084
#status preliminary: translated from GB/EMBL/DBJ
#molecule-type mRNA
#residues 1-207 ##label BRA
#cross-references GB:LI16914; NID:9438519; PIDN:AAA16620.1; PID:9438520
CLASSIFICATION #superfamily Interleukin-6
SUMMARY #length 207 #molecular-weight 23212 #checksum 9069

Query Match 10.0%; Score 158; DB 2; Length 207;
Best Local Similarity 28.2%; Pred. No. 3.83e-09;
Matches 29; Conservative 28; Mismatches 43; Indels 3; Gaps 2;

Db 30 GPLCGDATSNFLPTPA-DK--WEELIKYILGKISALKKEMCDNKKCEDSEALAEENL 86
      1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Oy 13 GSLVSGTRGRLPPAPPEFKDLQLRNNNAWVDECFRLCRGICRGILEPAIFL 72
      1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Db 87 NLPMIAEKDGCFOGSPFOETCIRITGLOEPDIYIKFLQDKY 129
      1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Oy 73 KLPAINDRHGCLIGFNETSCLKLADGFPEFVLFRTTFE 115
      1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1

RESULT 10
ENTRY #type complete
TITLE LRG1 protein - yeast (Saccharomyces cerevisiae)
ALTERNATE_NAMES protein D0764; protein YDL240w
ORGANISM #formal_name Saccharomyces cerevisiae
DATE 12-Jul-1996 #sequence_revision 12-Jul-1996 #text_change
07-May-1999
ACCESSIONS S67804; S43158; S47956
REFERENCE S67798
#authors Alt-Moerbe, J.; Schneider, C.; Moro, M.
#submission submitted to the Protein Sequence Database, July 1996
#accession S67804
##molecule-type DNA
##residues 1-1017 ##label ALT
#cross-references EMBL:Z72488; NID:91431407; PID:e253281; PID:91431408
MIPS:YDL240w
##experimental_source strain S288C
REFERENCE S43158
#authors Mueller, A.; Xu, G.; Wells, R.; Hollenberg, C.P.;
Piepersberg, W.
#submission submitted to the EMBL Data Library, March 1994
#description LRG1 is expressed during sporulation in Saccharomyces
cerevisiae and contains motifs homologous to LIM and
rho/racGAP domains.
#accession S43158
##molecule-type DNA
##residues 1-530, 'Q', 532-765, 'S', 767-790, 'T', 792-820, 'Q', 822-837,
'S', 839-848, 'L', 850-894, 'F', 896-927, 'T', 929-934, 'S',
936-976, 'TIL', 980, 'PPL', 995, 'KV', 998, 'OSIIPNVT',
#label MOE
#cross-references EMBL:X78453; NID:9468734; PID:9468735
S47956
#authors Mueller, L.; Xu, G.; Wells, R.; Hollenberg, C.P.;
Piepersberg, W.
#journal Nucleic Acids Res. (1994) 22:3151-3154
#title LRG1 is expressed during sporulation in Saccharomyces
cerevisiae and contains motifs similar to LIM and
rho/racGAP domains.
#cross-references MUID:94344779
#accession S47956
#molecule-type DNA
#residues 28-89:98-136, 'MF', 139-148:157-184:419-474:755-765, 'S',
767-790, 'T', 792-820, 'Q', 822-837, 'S', 839-848, 'L',
850-894, 'F', 896-910 ##label MD2
#cross-references EMBL:X78453

GENETICS
#gene SGD:LRG1
#cross-references SGD:S0002399; MIPS:YDL240w
#map_position 4L
CLASSIFICATION #superfamily LIM metal-binding repeat homology

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KEYWORDS      transmembrane protein
FEATURE
28-89         #domain LIM metal-binding repeat homology #label LIM1\
96-148        #domain LIM metal-binding repeat homology #label LIM2\
157-184       #domain LIM metal-binding repeat homology #status
348-364       #domain transmembrane #status predicted #label TM1\
419-474       #domain LIM metal-binding repeat homology #label LIM4\
467-503       #domain transmembrane #status predicted #label TM2\
SUMMARY
Query Match   7.1%; Score 112; DB 2; Length 1017;
Best Local Similarity 26.3%; Pred. No. 2,55e-02;
Matches 21; Conservative 23; Mismatches 33; Indels 3; Gaps 3;

Db 808 10LSALKKFIRELPDPISTDLVEMITKAARIDLEDEKQVILLIYSLPTVNRNLLER 867
:: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::
Oy 102 FEFVFLFKFLTEFGKSVINVDVMEILTKTLGMDIOEENKLTKTHTSP -PKFDRLGLR 160
:: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::

Db 868 LLSFLHWTSSP-S-YIENEM 885
:: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::
Oy 161 LOGLRVVRHFASFVYLSAM 180
:: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::

RESULT 11
ENTRY A42247 #type complete
TITLE myelomonocytic growth factor precursor - chicken
ALTERNATE_NAMES colony-stimulating factor CMGF
ORGANISM #formal_name Gallus gallus common_name chicken
DATE 16-Feb-1994 #sequence_revision 18-Nov-1994 #text_change
16-Jul-1989

ACCESSIONS A42247; S03633
REFERENCE A42247
#authors Sterneck, E.; Blattner, C.; Graf, T.; Leutz, A.
#journal Mol. Cell. Biol. (1992) 12:1728-1735
#title Structure of the chicken myelomonocytic growth factor gene
and specific activation of its promoter in avian
myelomonocytic cells by protein kinases.

#cross-references MUID:92195319
#accession A42247
#status preliminary
#molecule_type DNA
#residues 1-201 ##label STF
#note sequence extracted from NCBI backbone (NCBIN:89832,
NCBIP:89836)

REFERENCE S03633
#authors Leutz, A.; Damm, K.; Sterneck, E.; Kowenz, E.; Ness, S.;
Frank, R.; Gausepohl, H.; Pan, Y.C.E.; Smart, J.; Hayman,
M.; Graf, T.
#journal EMBO J. (1989) 8:175-181
#title Molecular cloning of the chicken myelomonocytic growth factor
(CMGF) reveals relationship to interleukin 6 and
granulocyte colony stimulating factor.

#cross-references MUID:89231616
#accession S03633
#molecule_type mRNA
#residues 1-201 ##label LEU
#cross-references EMBL:X14477; NID:963596; PIDD:CA32639.1; PID:963597
CLASSIFICATION #superfamily Interleukin-6
KEYWORDS glycoprotein

FEATURE
1-23 #domain signal sequence #status predicted #label SIG\
24-101 #product myelomonocytic growth factor #status predicted
#label MAT\
123,137 #binding site carbonylrate (Asn) (covalent) #status
predicted
SUMMARY #length 201 #molecular-weight 22373 #checksum 9000

Query Match 7.0%; Score 111; DB 2; Length 201;
Best Local Similarity 20.5%; Pred. No. 3.48e-02;
Matches 23; Conservative 32; Mismatches 53; Indels 4; Gaps 4;

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OY 81 DHGCLIGENESCCLKLADGFEFF-EVLFKFLTEFFGKSVINVDVWELLTITGLMDIOEE 139  
Matches 23; Conservative 14; Mismatches 20; Indels 7; Gaps 7;

Db 146 MEDICLDTVTLPADQRPPTFG-PF-QOOVGGFFIIANFORLEAYRAL 195  
OY 140 LNKLTHTYSPKFDRLGLRLOGIKYVWRHFAFSPYVLSAMEKFAQAVRVL 191

RESULT 12  
ENTRY F48563 #type complete  
TITLE I7 protein - fowlpox virus (strain HP444)  
ORGANISM #formal\_name fowlpox virus  
DATE 17-Feb-1994 #sequence\_revision 17-Feb-1994 #text\_change 16-Jul-1999

ACCESSIONS F48563  
REFERENCE A48563  
#authors Blinn, M.M.; Boursnell, M.E.; Skinner, M.A.  
#journal Virus Res. (1992) 24:161-172  
#title Gene translocations in poxviruses: the fowlpox virus thymidine kinase gene is flanked by 15 bp direct repeats and occupies the locus which in vaccinia virus is occupied by the ribonucleotide reductase large subunit gene.

#cross-references MUID:92410746  
#accession F48563  
#molecule\_type DNA  
#residues 1-421 #label BIN  
#cross-references GB:AJ223385; NID:g3123522; PIDN:CA111298.1; PID:e1292198; PID:g3123535  
#note sequence extracted from NCBI backbone (NCBI:113549, NCBI:113555)

GENETICS I7  
CLASSIFICATION #superfamily vaccinia virus I7 protein  
KEYWORDS late protein  
SUMMARY #length 421 #molecular-weight 48621 #checksum 8049

Query Match 6.9%; Score 109; DB 1; Length 421;  
Best Local Similarity 36.4%; Pred. No. 6.45e-02;  
Matches 16; Conservative 13; Mismatches 13; Indels 2; Gaps 2;

Db 282 GLNRNSYLSLANENADIDLFNFIDNYGYTAGCINVEVQQLT 325  
OY 87 GFNETSCCLKLADGFEFFEVLFKFLTEFFG-KSV-INVDVWELLT 128

RESULT 13  
ENTRY C70411 #type complete  
TITLE hypothetical protein ag\_1286 - Aquifex aeolicus  
ORGANISM #formal\_name Aquifex aeolicus  
DATE 08-May-1998 #sequence\_revision 08-May-1998 #text\_change 08-May-1998

ACCESSIONS C70411  
REFERENCE A70300  
#authors Decker, G.; Warren, P.V.; Gaasterland, T.; Young, W.G.; Lennox, A.L.; Graham, D.E.; Overbeek, R.; Shead, M.A.; Keller, M.; Aulay, M.; Huber, R.; Feldman, R.A.; Short, J.M.; Olson, G.O.; Swanson, R.V.  
#journal Nature (1998) 392:353-358  
#title The complete genome of the hyperthermophilic bacterium Aquifex aeolicus.

#cross-references MUID:98196666  
#accession C70411  
#status preliminary; nucleic acid sequence not shown; translation not shown

GENETICS #molecule\_type DNA  
#residues 1-281 #label AOF  
#cross-references GB:AE000732; NID:g2983704; PID:g2983716; GB:AE000657  
#experimental\_source strain VFS

GENETICS ag\_1286  
#gene #length 281 #molecular-weight 32624 #checksum 4784  
SUMMARY Query Match 6.8%; Score 108; DB 2; Length 281;

Best Local Similarity 35.9%; Pred. No. 8.75e-02;  
Matches 23; Conservative 14; Mismatches 20; Indels 7; Gaps 7;

Db 170 EGLEFFLV-FSKGKVINIAPLVLSLRDIDMELAKRI-ADYREKFPKQKDL-M 226  
OY 105 EVLFKFLTEFFGKSVINVDVWELLTK-TLGMIDQIEELNLTHTYSPK-F-D-RGLILGR 160

Db 227 VEGM 230  
OY 161 LQGL 164

RESULT 14  
ENTRY JC5495 #type complete  
TITLE Prox 1 protein - chicken  
ORGANISM #formal\_name Gallus gallus  
DATE 07-Jul-1997 #sequence\_revision 29-Aug-1997 #text\_change 17-Oct-1997

ACCESSIONS JC5495  
REFERENCE JC5495  
#authors Tomarev, S.I.; Sundin, O.; Banerjee-Basu, S.; Duncan, M.K.; Yang, J.M.; Platigorsky, J.  
#journal Dev. Dyn. (1996) 206:354-367  
#title Chicken homeobox gene Prox 1 related to Drosophila prospero is expressed in the developing lens and retina.

#cross-references MUID:97006692  
#contents len  
#accession JC5495  
#molecule\_type mRNA  
#residues 1-736 #label TOM  
#cross-references GB:U46563  
#note This protein is involved in eye development and function.

COMMENT #length 736 #molecular-weight 83086 #checksum 1607  
SUMMARY

Query Match 6.8%; Score 108; DB 2; Length 736;  
Best Local Similarity 41.7%; Pred. No. 8.75e-02;  
Matches 20; Conservative 11; Mismatches 12; Indels 5; Gaps 4;

Db 600 NML-KTFPSDYKFNRCITSQ--IK-WFSNPREYII-QMEKYARQAI 642  
OY 141 NKLTKHTYSPKFDRLGLRLOGIKYVWRHFAFSPYVLSAMEKFAQAV 188

RESULT 15  
ENTRY JE0269 #type complete  
TITLE Prox1 protein - mouse  
ORGANISM #formal\_name Mus musculus  
DATE 05-Feb-1999 #sequence\_revision 05-Feb-1999 #text\_change 17-Mar-1999

ACCESSIONS JE0269  
REFERENCE JE0269  
#authors Tomarev, S.I.; Zinovleva, R.D.; Chang, B.; Hawes, N.L.  
#journal Biochem. Biophys. Res. Commun. (1998) 248:684-689  
#title Characterization of the mouse Prox1 gene.

#cross-references MUID:98369610  
#accession JE0269  
#molecule\_type mRNA  
#residues 1-737 #label TOM  
#cross-references GB:AF061576

GENETICS #map\_position 1  
#introns 575/3; 611/3; 676/3  
SUMMARY #length 737 #molecular-weight 83126 #checksum 4650

Query Match 6.8%; Score 108; DB 2; Length 737;  
Best Local Similarity 41.7%; Pred. No. 8.75e-02;  
Matches 20; Conservative 11; Mismatches 12; Indels 5; Gaps 4;

Db 601 NML-KTFPSDYKFNRCITSQ--IK-WFSNPREYII-QMEKYARQAI 643  
OY 141 NKLTKHTYSPKFDRLGLRLOGIKYVWRHFAFSPYVLSAMEKFAQAV 188

Fri Sep 15 16:35:04 2000

US-09-230-048-2.rpt

Page 8

Search completed: Fri Sep 15 16:32:53 2000  
Job time : 31 secs.

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FT CARBOHYD 172 172 POTENTIAL.  
SQ SEQUENCE 212 AA; 23668 MW; C73C035226BA4B9F CRC64;  
Query Match 16.6%; Score 262; DB 1; Length 212;  
Best Local Similarity 28.4%; Pred. No. 2,95e-32;  
Matches 40; Conservative 36; Mismatches 65; Indels 0; Gaps 0;

DB 69 KETCRSNMCDSTKALNNLTPKMAKDCGFGFNEPCTCYKITGGLFEFVLEY 128  
OY 51 RDLCTRTGICGKLEPPAIFHLKLPAINDDHCGLIGFNETSCLKLDGFFFEVLEKF 110  
DB 129 LQNRSESEBARAVOMSTKVLIOLOKAKNLDAITTPPTNLSLTKLQAOQMLQD 188  
OY 111 LTTEFGKSVINVDWMLLTKLGMIOELNKLTKHTHSPKFRDGLGRLQGLKTVVRH 170  
DB 189 MTHILRSFEFLQSSLRAL 209  
OY 171 FASFYVLSAMEKFAQAVRVL 191

RESULT 2  
ID IL6\_MACFA STANDARD; PRT; 212 AA.  
AC P79341;  
DT 15-JUL-1998 (Rel. 36, Created)  
DT 15-JUL-1998 (Rel. 36, Last sequence update)  
DT 15-JUL-1998 (Rel. 36, Last annotation update)  
DE INTERLEUKIN-6 PRECURSOR (IL-6).  
CN IL6.  
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;  
OC Cercopithecoidea; Macaca.  
RN [1]  
RP SEQUENCE FROM N.A.  
RA Tatum M.;  
RL Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.  
CC -1- FUNCTION: IL6 IS A CYTOKINE WITH A WIDE VARIETY OF BIOLOGICAL  
FUNCTIONS: IT PLAYS AN ESSENTIAL ROLE IN THE FINAL DIFFERENTIATION  
OF B-CELLS INTO IG-SECRETING CELLS. IT INDUCES MYELOMA AND  
PLASMACYTOMA GROWTH. IT INDUCES NERVE CELLS DIFFERENTIATION, IN  
HEPATOCYTES IT INDUCES ACUTE PHASE REACTANTS (BY SIMILARITY).  
CC -1- SIMILARITY: BELONGS TO THE IL-6 SUPERFAMILY.  
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CC -----  
DR EMBL; AB000554; BAA19148.1; -  
DR HSSP; P05231; 21L6.  
DR PFAM; PF00489; IL6; 1.  
DR PRINTS; PR00433; IL6GCSFMGF.  
DR PROSITE; PS00434; INTERLEUKIN-6.  
DR PROSITE; PS00254; INTERLEUKIN-6; 1.  
KW Cytokine; Glycoprotein; Growth factor; Signal.  
FT SIGNAL 1 29  
FT CHAIN 1 29 BY SIMILARITY.  
FT DISULFID 72 78 INTERLEUKIN-6.  
FT DISULFID 101 111 POTENTIAL.  
FT CARBOHYD 73 73 POTENTIAL.  
FT CARBOHYD 172 172 POTENTIAL.  
SQ SEQUENCE 212 AA; 23654 MW; CR8173FCBF0B0389 CRC64;

Query Match 16.3%; Score 258; DB 1; Length 212;  
Best Local Similarity 28.4%; Pred. No. 2.10e-31;  
Matches 40; Conservative 36; Mismatches 65; Indels 0; Gaps 0;

DB 69 KETCRSNMCDSTKALNNLTPKMAKDCGFGFNEPCTCYKITGGLFEFVLEY 128  
OY 51 RDLCTRTGICGKLEPPAIFHLKLPAINDDHCGLIGFNETSCLKLDGFFFEVLEKF 110

DB 129 LQNRSESEBARAVOMSTKVLIOLOKAKNLDAITTPPTNLSLTKLQAOQMLQD 188  
OY 111 LTTEFGKSVINVDWMLLTKLGMIOELNKLTKHTHSPKFRDGLGRLQGLKTVVRH 170  
DB 189 MTHILRSFEFLQSSLRAL 209  
OY 171 FASFYVLSAMEKFAQAVRVL 191

RESULT 3  
ID IL6\_PIG STANDARD; PRT; 212 AA.  
AC P26893;  
DT 01-AUG-1992 (Rel. 23, Created)  
DT 01-AUG-1992 (Rel. 23, Last sequence update)  
DT 15-DEC-1998 (Rel. 37, Last annotation update)  
DE INTERLEUKIN-6 PRECURSOR (IL-6).  
CN IL6.  
OS Sus scrofa (Pig).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Sus.  
RN [1]  
RP SEQUENCE FROM N.A.  
RA MEDLINE; 91338547.  
RA Richards C., Saklatvala J.;  
RT "Molecular cloning and sequence of porcine interleukin 6 cDNA and  
expression of mRNA in synovial fibroblasts in vitro."  
RL Cytokine 3:269-276(1991).  
RN [2]  
RP SEQUENCE FROM N.A.  
RA MEDLINE; 92360284.  
RA Mathlagan N., Bixby J.A., Roberts M.R.;  
RT "Expression of interleukin-6 in porcine, ovine, and bovine  
preimplantation conceptuses."  
RL Mol. Reprod. Dev. 32:324-330(1992).  
CC -1- FUNCTION: IL6 IS A CYTOKINE WITH A WIDE VARIETY OF BIOLOGICAL  
FUNCTIONS: IT PLAYS AN ESSENTIAL ROLE IN THE FINAL DIFFERENTIATION  
OF B-CELLS INTO IG-SECRETING CELLS. IT INDUCES MYELOMA AND  
PLASMACYTOMA GROWTH. IT INDUCES NERVE CELLS DIFFERENTIATION, IN  
HEPATOCYTES IT INDUCES ACUTE PHASE REACTANTS.  
CC -1- SIMILARITY: BELONGS TO THE IL-6 SUPERFAMILY.  
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or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
CC -----  
DR EMBL; M86722; AAC37333.1; -  
DR HSSP; M80258; AAC27127.1; -  
DR PFAM; PF00489; IL6; 1.  
DR PRINTS; PR00433; IL6GCSFMGF.  
DR PROSITE; PS00434; INTERLEUKIN-6.  
DR PROSITE; PS00254; INTERLEUKIN-6; 1.  
KW Cytokine; Glycoprotein; Growth factor; Signal.  
FT SIGNAL 1 29  
FT CHAIN 1 29 BY SIMILARITY.  
FT DISULFID 72 78 INTERLEUKIN-6.  
FT DISULFID 101 111 BY SIMILARITY.  
FT CONFLICT 30 30 G -> E (IN REF. 2).  
SQ SEQUENCE 212 AA; 23880 MW; EF100ED030B6FD0 CRC64;

Query Match 16.3%; Score 257; DB 1; Length 212;  
Best Local Similarity 25.5%; Pred. No. 3.43e-31;  
Matches 36; Conservative 47; Mismatches 58; Indels 0; Gaps 0;

DB 69 KETCRSNMCDSTKALNNLTPKMAKDCGFGFNEPCTCYKITGGLFEFVLEY 128  
OY 51 RDLCTRTGICGKLEPPAIFHLKLPAINDDHCGLIGFNETSCLKLDGFFFEVLEKF 110

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Db 129 LOKESKNGKNGVAVQVISTKALIQTLROKGRKPPDATTNPNTNGLDKLOSONEMKN 188
Oy 111 LTTEGKSVINVDVWELLTKTLGMDIQEELNKLTKHTSPFNRGGLRGLQKTYWVR 170
Db 189 TKILIRSLDFLOFLSLRAI 209
Oy 171 FASFVLSAMEKFAAGAVRVL 191

RESULT 4
ID IL6 MACMU STANDARD: PRT: 212 AA.
AC P51494:
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE INTERLEUKIN-6 PRECURSOR (IL-6).
GN IL6.
OS Macaca mulatta (Rhesus macaque).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecinae; Macaca.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-RAC 2:
RX MEDLINE: 96003435.
RA Villinger F.O., Brar S.S., Wayne A.E., Chikala N., Ansari A.A.;
RT "Comparative sequence analysis of cytokine genes from human and
RT nonhuman primates.";
RL J. Immunol. 155:3946-3954(1995).
CC -1- FUNCTION: IL6 IS A CYTOKINE WITH A WIDE VARIETY OF BIOLOGICAL
CC ACTIONS: IT PLAYS AN ESSENTIAL ROLE IN THE FINAL DIFFERENTIATION
CC OF B-CELLS INTO IG-SECRETING CELLS, IT INDUCES MYELOMA AND
CC PLASMACYTOMA GROWTH, IT INDUCES NERVE CELLS DIFFERENTIATION, IN
CC HEPATOCYTES IT INDUCES ACUTE PHASE REACTANTS (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE IL-6 SUPERFAMILY.
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CC EMBL: L26028; AAA9978.1; -.
CC DR HSSP: P05231; 1ALU.
CC DR PFAM: PF00489; IL6.1.
CC DR PRINTS: PR00433; IL6GSPMGF.
CC DR PRINTS: PR00434; INTERLEUKIN6.
CC DR PROSITE: PS00254; INTERLEUKIN_6; 1.
CC KW Cytokine; Glycoprotein; Growth factor; Signal.
CC FT SIGNAL 1 29 BY SIMILARITY.
CC FT CHAIN 30 212 INTERLEUKIN-6.
CC FT DISULFID 72 78 POTENTIAL.
CC FT DISULFID 101 111 POTENTIAL.
CC FT CARBOHYD 73 73 POTENTIAL.
CC FT CARBOHYD 172 172 POTENTIAL.
CC SQ SEQUENCE 212 AA; 23728 MW; 4130DFE0CF0BCAD CRC64;

Query Match: 16.2%; Score 256; DB 1; Length 212;
Best Local Similarity 28.4%; Pred. No. 5,60e-31;
Matches 40; Conservative 35; Mismatches 66; Indels 0; Gaps 0;

Db 69 KETCRNSNCKSSKSLLENNLNLPMMAEKDCGFCGFMEDCLVYITIGLLEFVYLEY 128
Oy 51 RDLCTRTGCKCKILPRAALFHKLRAINDYDGCGLGFNEISGCKRLADGFFEFVLEF 110
Db 129 LONRESSEEOARAVOMSTKVLIOFLQKKAKMLDATTEPPTNMSLTKLOAONOMOD 188
Oy 111 LTTEGKSVINVDVWELLTKTLGMDIQEELNKLTKHTSPFNRGGLRGLQKTYWVR 170
Db 189 TKILIRSLDFLOFLSLRAI 209
Oy 171 FASFVLSAMEKFAAGAVRVL 191
Db 169 MTHLILMSFKFLOSINLRAI 209
Oy 111 LTTEGKSVINVDVWELLTKTLGMDIQEELNKLTKHTSPFNRGGLRGLQKTYWVR 170

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OY		171	EASFVYLASMEKFAAGAVRL	191
ID	RESULT	5	STANDARD;	PRT; 212 AA.
AC	IL6 HUMAN			
DT	P05231;			
DT	13-AUG-1987 (Rel. 05, Created)			
DT	13-AUG-1987 (Rel. 05, Last sequence update)			
DE	15-JUL-1998 (Rel. 36, Last annotation update)			
DE	INTERLEUKIN-6 PRECURSOR (IL-6) (B-CELL STIMULATORY FACTOR 2) (BSF-2)			
DN	(INTERFERON BETA-2) (HYBRIDOMA GROWTH FACTOR).			
GN	IL6 OR IFNB2.			
OS	Homo sapiens (Human).			
OC	Eumetazoa: Chordata: Craniata: Vertebrata; Euteleostomi;			
CC	Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.			
RN	[1]			
RP	SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.			
RX	MEDLINE: 87065033.			
RA	Hirano T., Yasukawa K., Harada H., Taga T., Watanabe Y., Matsuda T.,			
RA	Kashiyamamura S.-i., Nakajima K., Koyama K., Iwamoto A., Tsunatsawa S.,			
RA	Sakigaya F., Matsuji H., Takahara Y., Taniguchi T., Kishimoto T.;			
RT	"Complementary DNA for a novel human interleukin (BSF-2) that induces			
RT	B lymphocytes to produce immunoglobulin."			
RL	Nature 324:73-76(1986).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE: 88082664.			
RA	Yasukawa K., Hirano T., Watanabe Y., Muratani K., Matsuda T.,			
RA	Nakai S., Kishimoto T.;			
RT	"Structure and expression of human B cell stimulatory factor-2			
RT	(BSF-2/IL-6) gene."			
RL	EMBO J. 6:2939-2945(1987).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE: 87067433.			
RA	May L.T., Helfgott D.C., Sehgal P.B.;			
RT	"Anti-beta-interferon antibodies inhibit the increased expression of			
RT	HLA-B7 mRNA in tumor necrosis factor-treated human fibroblasts:			
RT	structural studies of the beta 2 interferon involved."			
RL	Proc. Natl. Acad. Sci. U.S.A. 83:8957-8961(1986).			
RN	[4]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE: 87053818.			
RA	Zilberstein A., Ruggieri R., Korn J.H., Revel M.;			
RT	"Structure and expression of cDNA and genes for human			
RT	interferon-beta-2, a distinct species inducible by growth-stimulatory			
RT	cytokines."			
RL	EMBO J. 5:2529-2537(1986).			
RN	[5]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE: 88088768.			
RA	Brakenhoff J.P.J., de Groot E.R., Evers R.F., Pannekoek H.,			
RA	Aarden L.A.;			
RT	"Molecular cloning and expression of hybridoma growth factor in			
RT	Escherichia coli."			
RL	J. Immunol. 139:4116-4121(1987).			
RN	[6]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE: 89391958.			
RA	Tonouchi N., Miya K., Karasuyama H., Matsui H.;			
RT	"Deletion of 3' untranslated region of human BSF-2 mRNA causes			
RT	stabilization of the mRNA and high-level expression in mouse NIH3T3			
RT	cells."			
RL	Biochem. Biophys. Res. Commun. 163:1056-1062(1989).			
RN	[7]			
RP	SEQUENCE FROM N.A.			
RX	TISSUE-FIBROBLAST;			
RC	MEDLINE: 87004683.			
RA	Haegeman G., Content J., Volckaert G., Derynck R., Tavernier J.,			
RA	Fiers W.;			
RT	"Structural analysis of the sequence coding for an inducible 26-kDa			
RT	protein in human fibroblasts."			
RL	Eur. J. Biochem. 159:625-632(1986).			

RA SEQUENCE FROM N.A.  
 RX MEDLINE: 89193317.  
 RA Wong G., Witek-Giamotti J., Hewick R., Clark S., Ogawa M.;  
 RA "Interleukin 6: Identification as a hematopoietic colony-stimulating  
 RT factor.";  
 RL Behring Inst. Mitt. 83:40-47(1988).  
 RN [9]  
 RN SEQUENCE FROM N.A.  
 RX MEDLINE: 93178270.  
 RA Chen Q.Y.;  
 RT "Stable and efficient expression of human interleukin-6 cDNA in  
 RT mammalian cells after gene transfer.";  
 RL Chang-Hua Chung Liu Tse Chih 14:340-344(1992).  
 RN [10]  
 RP SEQUENCE OF 30-63.  
 RX MEDLINE: 88154445.  
 RA van Damme J., van Beunnen J., Decock B., van Snick J., de Ley M.,  
 RA Billiau A.;  
 RT "Separation and comparison of two monokines with  
 RT lymphocyte-activating factor activity: IL-1 beta and hybridoma growth  
 RT factor (HGF). Identification of leukocyte-derived HGF as IL-6.";  
 RL J. Immunol. 140:1534-1541(1988).  
 RN [11]  
 RP SEQUENCE OF 50-212 OF RECOMBINANT FORM LACKING 1ST DISULFIDE BOND.  
 RX MEDLINE: 95154344.  
 RA Bleton J., la Flura A., Bertolero F., Orsini G., Valasina B.,  
 RA Ziliocto R., de Filippis V., Polyverino de Laureto P., Fontana A.;  
 RT "Structure, stability and biological properties of a N-terminally  
 RT truncated form of recombinant human interleukin-6 containing a single  
 RT disulfide bond.";  
 RL Eur. J. Biochem. 227:573-581(1995).  
 RN [12]  
 RP DISULFIDE BONDS.  
 RX MEDLINE: 89286115.  
 RA Clogston C.L., Boone T.C., Grandall B.C., Mendiaz E.A., Lu H.S.;  
 RT "Disulfide structures of human interleukin-6 are smaller to those of  
 RT human granulocyte colony stimulating factor.";  
 RL Arch. Biochem. Biophys. 272:144-151(1989).  
 RN [13]  
 RN MUTAGENESIS.  
 RP MEDLINE: 91243808.  
 RX Lueticken C., Krueitgen A., Moeller C., Helnrich P.C., Rose-John S.;  
 RT "Evidence for the importance of a positive charge and an  
 RT alpha-helical structure of the C-terminus for biological activity of  
 RT human IL-6.";  
 RL FEBS Lett. 282:265-267(1991).  
 RN [14]  
 RP STRUCTURE BY NMR.  
 RX MEDLINE: 96134845.  
 RA Nishimura C., Matanabe A., Gouda H., Shimada I., Arata Y.;  
 RT "Folding topologies of human interleukin-6 and its mutants as studied  
 RT by NMR spectroscopy.";  
 RL Biochemistry 35:273-281(1996).  
 RN [15]  
 RP STRUCTURE BY NMR.  
 RX MEDLINE: 97303053.  
 RA Xu G.-Y., Yu H.-A., Hong J., Stahl M., McDonagh T., Kay L.E.,  
 RA Cumming D.A.;  
 RT "Solution structure of recombinant human interleukin-6.";  
 RL J. Mol. Biol. 268:468-481(1997).  
 RN [16]  
 RP X-RAY CRYSTALLOGRAPHY (1.9 ANGSTROMS).  
 RX MEDLINE: 97224126.  
 RA Somers W., Stahl M., Seehra J.S.;  
 RT "1.9-A crystal structure of interleukin 6: implications for a novel  
 RT mode of receptor dimerization and signaling.";  
 RL EMBO J. 16:989-997(1997).  
 CC -1- FUNCTION: IL6 IS A CYTOKINE WITH A WIDE VARIETY OF BIOLOGICAL  
 CC OF B-CELLS INTO IG-SECRETING CELLS. IT INDUCES MYELOMA AND  
 CC PLASMACYTOMA GROWTH. IT INDUCES NERVE CELLS DIFFERENTIATION, IN  
 CC HEPATOCYTES IT INDUCES ACUTE PHASE REACTANTS.

```

CC -1 SIMILARITY: BELONGS TO THE IL-6 SUPERFAMILY.
CC -----
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CC -----
CC EMBL: X04430; CAA28026.1; -
CC DR EMBL: M14584; AAA52728.1; -
CC DR EMBL: X04602; CAA28268.1; -
CC DR EMBL: Y00081; CAA68278.1; -
CC DR EMBL: M18403; AAA52729.1; -
CC DR EMBL: M29150; AAA59154.1; -
CC DR EMBL: X04402; CAA27990.1; -
CC DR EMBL: X04403; CAA27991.1; -
CC DR EMBL: M54894; AAC41704.1; -
CC DR EMBL: S56892; AAD13886.1; -
CC DR EMBL: A09363; CAA00839.1; -
CC DR PIR: A32648; IVH0B2.
CC DR PIR: A25921; A25921.
CC DR PDB: 1IL6; 04-FEB-98.
CC DR PDB: 2IL6; 04-FEB-98.
CC DR PDB: 1ALU; 03-JUN-98.
CC DR MIM: 147620; -
CC DR PFAM: PF00489; IL6; 1.
CC DR PRINTS: PRO0433; IL6CSFMF.
CC DR PRINTS: PRO0434; INTERLEUKIN6.
CC DR PROSITE: PS00254; INTERLEUKIN_6; 1.
CC DR Cycokine; Glycoprotein; Growth factor; Signal; 3D-structure.
CC KM SIGNAL 1 29
CC FT CHAIN 30 212 INTERLEUKIN-6.
CC FT DISULFID 72 78
CC FT DISULFID 101 111
CC FT CARBOHYD 73 73
CC FT MUTAGEN 173 173 A->Y: ALMOST NO LOSS OF ACTIVITY.
CC FT MUTAGEN 185 185 W->R: NO LOSS OF ACTIVITY.
CC FT MUTAGEN 204 204 S->D: 13% ACTIVITY.
CC FT MUTAGEN 210 210 R->K,E,Q,T,A,P: LOSS OF ACTIVITY.
CC FT MUTAGEN 212 212 M->T,N,S,R: LOSS OF ACTIVITY.
CC SQ SEQUENCE 212 AA; 23718 MM; 1IED1FE1B724079 CRC64;

Query Match 16.0%; Score 257; DB 1; Length 212;
Best Local Similarity 27.7%; Pred. No. 3,96e-30;
Matches 39; Conservative 37; Mismatches 65; Indels 0; Gaps 0;

Db 69 KETCKRNNKCESSEKRALNNLNLPKMAEKDCQFSGNEEFCAVKITITGLFEFVLEY 128
QY ::::: ::::: ::::: ::::: ::::: ::::: ::::: ::::: ::::: :::::
51 ROLCYRTGICGKLEPAIPFIHKLPAINDIDHCGIIGENISCLKLKDGFEEFVLRKF 110
Db 129 LONRESSEBOARAVOMSTKYLIOFLQKAKNLAITTPDPTNSSLTKLOAONQIOD 188
QY 111 LTTEGKASIVNDVVELLTKTLGMDIQELNKLTRTHYSPKFRDLGLRIGLQKLVYRH 170
Db 189 MTHILRSFKEFLQSSLRAL 209
QY ::::: ::::: ::::: ::::: ::::: ::::: ::::: ::::: :::::
171 FASFYVLSAMEKFAQAVAYVL 191

RESULT 6
AC 095181; 019007; 046568; STANDARD: PRT: 208 AA.
DT 01-NOV-1987 (Rel. 35, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE INTERLEUKIN-6 PRECURSOR (IL-6).
GN IL6.
OS Equus caballus (Horse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Equidae; Equus.
NC [1]

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CC -----
DR EMBL; X57317; CAA40572.1; -.
DR PIR; S22162; S22162.
DR HSSP; P05231; 2IL6.
DR PFAM; PF00489; IL6; 1.
DR PRINTS; PR00433; IL6GSGFMG.
DR PROSITE; PS00254; INTERLEUKIN_6.
DR PROSITE; PS00254; INTERLEUKIN_6; 1.
DR CYTOKINE; Glycoprotein; Growth factor; Signal.
FT SIGNAL 1 29
FT CHAIN 30 208 BY SIMILARITY.
FT DISULFID 72 78 INTERLEUKIN-6.
FT DISULFID 101 111 BY SIMILARITY.
FT CARBOHYD 38 38 POTENTIAL.
SO SEQUENCE 208 AA; 23758 MW; A0F000B9A2EC341 CRC64;

Query Match 12.8%; Score 202; DB 1; Length 208;
Best Local Similarity 23.2%; Pred. No. 8,60e-20;
Matches 33; Conservative 48; Mismatches 55; Indels 6; Gaps 6;

Db 69 KEICRNDKDESSKKTLEAKNLKPKMEKDCFGSGFNOAICLTIRTAGLEVOYLYDY 128
OY 51 RDLCYRTGICGILPEPAIFHLKLPAINDDHCGLIGFNETSCCLKLADGFEFEVLEKF 110
DB 129 IQNVE-GNQE-NVRDL-RKNI-RTLIQLKOKIADLTTPAT-NTDLEKQSSNEWK 183
OY 111 LTTEFGKSVINVDVMEILLTKTLGMDIOBELN-KLTKRTHYSPKFRGLGRLGKLYVRH 169
DB 184 NAKILILNLNENLFOFSRLAI 205
OY 170 HFASFYVLSAMEKFAAGAVRL 191

RESULT 12
ID IL6_PROVI STANDARD: PRT; 209 AA.
AC Q28819;
DT 15-FEB-2000 (Rel. 39, Created)
DT 15-FEB-2000 (Rel. 39, Last sequence update)
DT 15-FEB-2000 (Rel. 39, Last annotation update)
DE INTERLEUKIN-6 PRECURSOR (IL-6) (FRAGMENT).
GN IL6.
OS Phoca vitulina (Harbor seal).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Carnivora; Plinipedidae; Phocidae; Phoca.
RN [1]
RN SEQUENCE FROM N.A.
RX MEDLINE; 96163018.
RX King D.P., Schrenzel M.D., McKnight M.L., Reiderson T.H., Hanni K.D.,
RA Stolt J.L., Ferrick D.A.;
RT "Molecular cloning and sequencing of interleukin 6 cDNA fragments from
RT the harbor seal (Phoca vitulina), killer whale (Orcinus orca), and
RT Southern sea otter (Enhydra lutris nereis).";
RL Immunogenetics 43:190-195(1996).
CC -1- FUNCTION: IL6 IS A CYTOKINE WITH A WIDE VARIETY OF BIOLOGICAL
CC FUNCTIONS: IT PLAYS AN ESSENTIAL ROLE IN THE FINAL DIFFERENTIATION
CC OF B-CELLS INTO IG-SECRETING CELLS, IT INDUCES MYELOMA AND
CC PLASMACYTOMA GROWTH, IT INDUCES NERVE CELLS DIFFERENTIATION, IN
CC HEPATOCYTES IT INDUCES ACUTE PHASE REACTANTS (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: SECRETED.
CC -1- SIMILARITY: BELONGS TO THE IL-6 SUPERFAMILY.
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CC -----
DR EMBL; L46802; AAB01430.1; -.
DR HSSP; P05231; 2IL6.
DR PFAM; PF00489; IL6; 1.

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DR PROSITE; PS00254; INTERLEUKIN_6; 1.
KW CYTOKINE; Glycoprotein; Growth factor; Signal.
FT SIGNAL 1 26
FT NON_TER 1
FT CHAIN 27 209 BY SIMILARITY.
FT DISULFID 69 75 INTERLEUKIN-6.
FT DISULFID 98 108 BY SIMILARITY.
SO SEQUENCE 209 AA; 23483 MW; 75144922E4B48E9 CRC64;

Query Match 12.7%; Score 201; DB 1; Length 209;
Best Local Similarity 22.6%; Pred. No. 1.36e-19;
Matches 30; Conservative 42; Mismatches 61; Indels 0; Gaps 0;

Db 66 KEMDKNKCKDSSEALAEENLRPKLAENKDCFGSGFNOETCLRTTGLERQIHLKY 125
OY 51 RDLCYRTGICGILPEPAIFHLKLPAINDDHCGLIGFNETSCCLKLADGFEFEVLEKF 110
DB 126 IQAVEGNKEDANSVYSTKLVLQMLKKVKSODEVTPPTPTDTSLOAILKAODKWLKH 185
OY 111 LTTEFGKSVINVDVMEILLTKTLGMDIOBELNKLTKRTHYSPKFRGLGRLGKLYVRH 170
DB 186 TTHILIRSLDF 198
OY 171 HFASFYVLSAMEK 183

RESULT 13
ID IL6_MARMO STANDARD: PRT; 207 AA.
AC O35736;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE INTERLEUKIN-6 PRECURSOR (IL-6).
GN IL6.
OS Marmota monax (Woodchuck).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Sciurinae;
OC Marmota.
RN [1]
RN SEQUENCE FROM N.A.
RX TISSUE-PERIPHERAL BLOOD;
RX MEDLINE; 98139533.
RX Lohrengel B., Lu M., Rogendorf M.;
RT "Molecular cloning of the woodchuck cytokines: TNF-alpha, IFN-gamma,
RT and IL-6.";
RL Immunogenetics 47:332-335(1998).
CC -1- FUNCTION: IL6 IS A CYTOKINE WITH A WIDE VARIETY OF BIOLOGICAL
CC FUNCTIONS: IT PLAYS AN ESSENTIAL ROLE IN THE FINAL DIFFERENTIATION
CC OF B-CELLS INTO IG-SECRETING CELLS, IT INDUCES MYELOMA AND
CC PLASMACYTOMA GROWTH, IT INDUCES NERVE CELLS DIFFERENTIATION, IN
CC HEPATOCYTES IT INDUCES ACUTE PHASE REACTANTS (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE IL-6 SUPERFAMILY.
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CC -----
DR EMBL; Y14139; CAA74571.1; -.
DR PFAM; PF00489; IL6; 1.
DR PRINTS; PR00433; IL6GSGFMG.
DR PRINTS; PR00434; INTERLEUKIN_6.
DR PROSITE; PS00254; INTERLEUKIN_6; 1.
KW CYTOKINE; Glycoprotein; Growth factor; Signal.
FT SIGNAL 1 18
FT CHAIN 19 207 BY SIMILARITY.
FT DISULFID 65 71 INTERLEUKIN-6.
FT DISULFID 94 104 POTENTIAL.
SO SEQUENCE 207 AA; 23770 MW; F30D19F86AD6A600 CRC64;

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FT    CONFLICT    173    187    AKLOSQSEEMLRHHTI -> LSCSHRRVAEAMNN (IN
FT    CONFLICT    200    201    FS -> LR (IN REF. 2).
SQ    SEQUENCE    208 AA; 23401 MW; 93B4456B2989CACC64;

Query Match          12.0%; Score 189; DB 1; Length 208;
Best Local Similarity 25.6%; Pred. No. 3, 3le-17;
Matches 3; Conservative 3; Mismatches 68; Indels 0; Gaps 0;

Db    65 KEMCDNRYKCEDESKAEALAEENLNLPKLAECGCEFCGSGENOETCLRTITVYGOEFOIYIKF 124
Oy    51 RDLICRTICGIGILEPAIFHKLPAINDPDHCGIGIGENHSCCLKLADGFFEEVYLKRF 110
Db    125 LODKESGEENKSVYTSFNVLLOWLKKRKNODEVTIPVPTVEVGLQAKLOSQSEEWLR 184
Oy    111 LITTEGKSVINVDVWELLTKTLGMDIOELNKLTKTHTSPKFDGLGRLQGLTWVRH 170
Db    185 TTIHTLRLEDF 197
Oy    171 FASFYVLSAMEKF 183

RESULT 15 STANDARD; PRT: 211 AA.
AC P20607;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-FEB-1991 (Rel. 17, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE INTERLEUKIN-6 PRECURSOR (IL-6).
GN IL6 OR IL-6.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]
RX MEDLINE; 89380206.
RA Northmann W., Blaciak T.A., Hattori M., Lee F., Fey G.H.;
RT "Structure of the rat interleukin 6 gene and its expression in
RT macrophage-derived cells."
RL J. Biol. Chem. 264:16072-16082(1989).
CC -1- FUNCTION: IL6 IS A CYTOKINE WITH A WIDE VARIETY OF BIOLOGICAL
CC FUNCTIONS: IT PLAYS AN ESSENTIAL ROLE IN THE FINAL DIFFERENTIATION
CC OF B-CELLS INTO IG-SECRETING CELLS, IT INDUCES MYELOMA AND
CC PLASMACYTOMA GROWTH, IT INDUCES NERVE CELLS DIFFERENTIATION, IN
CC HEPATOCYTES IT INDUCES ACUTE PHASE REACTANTS.
CC -1- SIMILARITY: BELONGS TO THE IL-6 SUPERFAMILY.
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CC -----
DR EMBL; M26744; AAA77659.1; -
DR EMBL; M26745; AAA41430.1; -
DR PIR; A34247; A34247.
DR HSSP; P05231; 1AUU.
DR PFAM; PF00489; IL6; 1.
DR PRINTS; PRO0433; IL6GCSFNGF.
DR PRINTS; PRO0434; INTERLEUKIN6.
DR PROSITE; PS00254; INTERLEUKIN_6; 1.
KW Cytokine; Growth factor; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 211
FT DISULFID 70 76
FT DISULFID 99 109
SQ SEQUENCE 211 AA; 24357 MW; 17D248A14FE96B5C1 CAC64;

Query Match          12.0%; Score 189; DB 1; Length 211;
Best Local Similarity 25.7%; Pred. No. 3, 3le-17;
Matches 3; Conservative 4; Mismatches 72; Indels 1; Gaps 1;

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Fri Sep 15 16:35:04 2000

US-09-230-048-2.rsp

Page 10

Db	55	ITYLREIETLEKREKELGNSGNSDALSNNKLPKEIQIKNOCGFGTGNOCJCLIKC	114
Qy	39	LNNMLWVIDEDEFRLDCTRTGICIKILEPALFHLKLAINDTBDGCLIGENNESCLKLA	98
Db	115	SGLLEFRYELFEYKNNLNODNKKDRAVYOSTETELVIAIFKOEIKDSYKILPTPTSNALL	174
Qy	93	DGEFEFEVLEFFLTERBKSYIAND-VNELLTKLUGDIOBELNKLKTRHYSPPKPDRLG	157
Db	175	MEKLESQEWLTKTIQILKALEEPLKVTMR	206
Qy	158	LGRIGLKYVWVRHFAFSFYVLSAMEKFGQAVR	189

Search completed: Fri Sep 15 16:33:31 2000  
Job time : 20 secs.







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AC 028403:
DT 01-NOV-1996 (Tremblrel. 01, Created)
DT 01-NOV-1996 (Tremblrel. 01, Last sequence update)
DT 01-NOV-1999 (Tremblrel. 12, Last annotation update)
DE INTERLEUKIN 6 (FRAGMENT).
GN IL-6.
OS Euhadra lutris (Sea otter).
OC Eukaryota; Metazoa; Chordata; Craniala; Vertebrata; Mammalia;
OC Eutheria; Carnivora; Fissipedia; Mustelidae; Euhadra.
RN [1]
RP SEQUENCE FROM N.A.
RA MEDLINE; 96163018.
RA KING D.P., SCHRENZEL M.D., MCKNIGHT M.L., REIDARSON T.H., HANNI K.D.,
RA STOTT J.L., FERRICK D.A.;
RT "Molecular cloning and sequencing of interleukin 6 cDNA fragments from
RT the harbor seal (Phoca vitulina), Killer whale (Orcinus orca), and
RT Southern sea otter (Euhadra lutris nereis).";
RL Immunogenetics 43:190-195(1996).
DR EMBL; I46804; AAB01428.1; -
DR HSSP; P05231; 21L6.
DR PROSITE; PS00254; INTERLEUKIN_6; 1.
DR PFAM; PF00489; IL-6; 1.
FT NON_TER
SQ SEQUENCE 207 AA; 23527 MW; 0F2CAC62 CRC32;

Query Match
Best local similarity 23.6%; Score 169; DB 6; Length 207;
Matches 33; Conservative 40; Mismatches 65; Indels 2; Gaps 2;

DB 67 EMCDDKYNKCEDSKVLAENNLPRLEAKDRCFQSFNOETCLRTITGLOEFOIHLKYL 126
QY 52 DLGRTGICGILBPAIFHLKLPAINDTDHCGLIGNETSCLMKADGFEEFVLEFRL 111
DB 127 ESNTEGKNDNAHYIISIKHL-LQTLRPMNOIEVTT-PDPTTDSIALFSSODKWLKHT 184
QY 112 TTEGKSYINVDVVELLTKTLGMDIQEELNKLTKTHYSPKFDGLGLRLOGLYKWWHF 171
DB 185 TIHLIRLEDFLOFSLRAI 204
QY 172 ASFYLSAMEKFAQAVRVL 191

RESULT 9
ID 09WV08 PRELIMINARY; PRT; 210 AA.
AC 09WV08:
DT 01-NOV-1999 (Tremblrel. 13, Created)
DT 01-NOV-1999 (Tremblrel. 12, Last sequence update)
DT 01-NOV-1999 (Tremblrel. 12, Last annotation update)
DE IL-6 (FRAGMENT).
OS Mesocricetus auratus (Golden hamster).
OC Eukaryota; Metazoa; Chordata; Craniala; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae; Mesocricetus.
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN-APA; TISSUE-KIDNEY;
RA NISHIDA E.;
RT "APA hamsters IL-6 partial cDNA";
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB028635; BAA78766.1; -
DR PROSITE; PS00254; INTERLEUKIN_6; 1.
FT NON_TER
SQ SEQUENCE 210 AA; 24060 MW; 0307F113 CRC32;

Query Match
Best local similarity 24.0%; Score 169; DB 11; Length 210;
Matches 40; Conservative 41; Mismatches 84; Indels 2; Gaps 2;

DB 39 PNRPVYTSQVGVGLTVLREIYELRKELCNMNGCDMDNDYVLENNLEPVQINDGC 98
QY 25 PDAREFEDLLIQR-LNMMLVVIDCFRDLCTYRGICIGLEPAIFHLKLPALINDDHC 83
DB 99 LQTSYWEICLLKTKTSGLDYQYLEFVYNNVODNKKARVIOSTIKTISQIFKQEVKG 158

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QY 84 GLIGFNETSCLKLADGFEFEVLEFKFLTTEFGKSYINVD-VNELTKTLGMDIQEELNK 142
DB 159 PDKVTYSPSTSKAILMAKESOKEMPTTKIKLKLAEFLEVTMR 205
QY 143 LTKTHYSPKFDRLGLRLOGLYKWWHFASFYLSAMEKFAQAVR 169

RESULT 10
ID 055041 PRELIMINARY; PRT; 101 AA.
AC 055041:
DT 01-JUN-1998 (Tremblrel. 06, Created)
DT 01-JUN-1998 (Tremblrel. 06, Last sequence update)
DT 01-NOV-1999 (Tremblrel. 12, Last annotation update)
DE INTERLEUKIN 6 (FRAGMENT).
GN IL-6.
OS Cricetusulus griseus (Chinese hamster).
OC Eukaryota; Metazoa; Chordata; Craniala; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae; Crictulus.
RN [1]
RP SEQUENCE FROM N.A.
RA HEINE H., DELUDE R.D., MONKS B., GOLENOCK D.T.;
RL Submitted (JAN-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF044667; AAC02100.1; -
DR HSSP; P05231; 21L6.
DR PROSITE; PS00254; INTERLEUKIN_6; 1.
DR PFAM; PF00489; IL-6; 1.
FT NON_TER
SQ SEQUENCE 101 AA; 11749 MW; DIDA362A CRC32;

Query Match
Best local similarity 25.5%; Score 121; DB 11; Length 101;
Matches 24; Conservative 27; Mismatches 42; Indels 1; Gaps 1;

DB 2 IORNDGCGYGYWEICLTKTSGLDYQYLEFVYNNVODNKKARVIOSTIKTISQI 61
QY 77 INDVHCGILGILNETSCLKLADGFEFEVLEFKFLTTEFGKSYINVD-VNELTKTLGMD 135
DB 62 FKQEVKDPDKRYMPSPTSKAILIEKLESOKWMP 95
QY 136 IQEELNKLTKTHYSPKFDRLGLRLOGLYKWWHF 169

RESULT 11
ID 046980 PRELIMINARY; PRT; 377 AA.
AC 046980:
DT 01-JUN-1998 (Tremblrel. 06, Created)
DT 01-JUN-1998 (Tremblrel. 06, Last sequence update)
DT 01-NOV-1999 (Tremblrel. 12, Last annotation update)
DE RIBULOSE BISPHOSPHATE CARBOXYLASE LARGE CHAIN (EC 4.1.1.39)
DE (FRAGMENT).
GN RBCL.
OS Synura uvelia.
OS Chloroplast.
OC Eukaryota; stramenopiles; Chrysophyceae; Synurales; Synura.
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN-CCM870;
RA DAVGUBERG N., ANDERSEN R.A.;
RL J. Phycol. 33:1031-1041(1997).
RC -1- FUNCTION: RUBISCO CATALYSES TWO REACTIONS: THE CARBOXYLATION OF D-
RIBULOSE 1,5-BISPHOSPHATE, THE PRIMARY EVENT IN PHOTOSYNTHETIC
CARBON DIOXIDE FIXATION, AS WELL AS THE OXIDATIVE FRAGMENTATION OF
THE PENTOSE SUBSTRATE IN THE PHOTORESPIRATION PROCESS. BOTH
REACTIONS OCCUR SIMULTANEOUSLY AND IN COMPETITION AT THE SAME
ACTIVE SITE.
CC -1- CATALYTIC ACTIVITY: D-RIBULOSE 1,5-BISPHOSPHATE + CO(2) -> 3-
PHOSHO-D-GLYCERATE.
CC -1- CATALYTIC ACTIVITY: D-RIBULOSE 1,5-BISPHOSPHATE + O(2) -> 3-
PHOSHO-D-GLYCERATE + 2-PHOSPHOGLYCOLATE.
CC -1- SUBUNIT: 8 LARGE CHAINS + 8 SMALL CHAINS.
DR EMBL; AF015586; AAC02925.1; -
DR PROSITE; PS00157; RUBISCO_LARGE; 1.

```

[illegible]





\*\*\*\*\*STN Columbus\*\*\*\*\*

FILE HOME ENTERED AT 16:30:20 ON 15 SEP 2000

=> file medicine

COST IN U.S. DOLLARS	ENTRY	SESSION	TOTAL
FULL ESTIMATED COST	0.15	0.15	

FILE MEDLINE ENTERED AT 16:30:25 ON 15 SEP 2000

FILE LAST UPDATED: 7 SEP 2000 (20000907/UP). FILE COVERS 1960 TO DATE.

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The OLD MEDLINE file segment now contains data from 1958 through 1965. Enter HELP CONTENT for details.

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=> s herpes virus type 8/ab,bi

33818 HERPES/BI  
294323 VIRUS/BI  
529956 TYPE/BI  
749460 8/BI  
5408175 AB/FA  
9 HERPES VIRUS TYPE 8/AB  
(HERPES(V)VIRUS(V)TYPE(V)8)BI (L) AB/FA)

33818 HERPES/BI  
294323 VIRUS/BI  
529956 TYPE/BI  
749460 8/BI

14 HERPES VIRUS TYPE 8/BI  
(HERPES(V)VIRUS(V)TYPE(V)8)BI  
14 HERPES VIRUS TYPE 8/AB,BI

=> s ll and interleukin#/ab,bi

90758 INTERLEUKIN#/BI  
5408175 AB/FA

59796 INTERLEUKIN#/AB  
(INTERLEUKIN#/BI (L) AB/FA)  
90758 INTERLEUKIN#/BI  
L2 1 LI AND INTERLEUKIN#/AB,BI

=> d bib ab

L2 ANSWER 1 OF 1 MEDLINE  
AN 97054369 MEDLINE  
DN 97054369  
TI The natural history and molecular heterogeneity of HIV-associated primary malignant lymphomatous effusions.  
AU Komanduri K V; Luce J A; McGrath M S; Herndler B G; Ng V L  
CS Department of Medicine, University of California, San Francisco, USA  
NC RO1 CA54742 (NCI)  
SO JOURNAL OF ACQUIRED IMMUNE DEFICIENCY SYNDROMES AND HUMAN RETROVIROLOGY.  
(1996 Nov 1) 13 (3) 215-26  
Journal code: B7J. ISSN: 1077-9450.  
CY United States

DT Journal Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 199702  
AB Primary malignant lymphomatous effusions arising in individuals infected with the human immunodeficiency virus, type 1 (HIV-1) represent a rare subset of HIV-associated lymphomas. Previous studies have demonstrated that the malignant cells are monoclonal (as defined by rearrangement of the immunoglobulin gene), express cell surface CD38, and are infected with Epstein-Barr virus (EBV) and human herpesvirus-8 (HHV-8). Despite these detailed molecular and immunophenotypic studies, clinical information on this disease entity is scant, prompting us to review the clinical features of eight cases seen at our institutions. All eight patients had total peripheral CD4+ lymphocytes < 200/microliter and presented with complaints related to body cavity distension. Routine laboratory values were nondiagnostic and yielded no prognostic information. Only two patients could tolerate and thus received chemotherapy with no obvious impact on their clinical course. The mean overall survival after diagnosis was 60 days (range 6-166 days).

Four patients were examined at autopsy. The primary malignant lymphomatous effusion either was the immediate cause of death or contributed significantly to the death of only two. All four patients examined post mortem, however, had lymphomatous infiltration of serosal surfaces adjacent to the site of the primary malignant effusion. Molecular immunologic studies performed on the malignant cells and effusion fluids revealed universal expression of cell surface CD38 and the presence of HHV-8 gene sequences, but in contrast with previous studies, only four had rearranged immunoglobulin genes or EBV present. IL-6 and IL-10 levels in the malignant effusion fluids were markedly elevated. In summary, this rare subset of HIV-associated lymphomas in our eight patients arose late in the course of HIV-associated disease, had a rapid clinical course, and was molecularly heterogeneous. A pathogenetic role for HHV-8 alone in this disease process is strengthened by our observation of four cases lacking EBV but containing HHV-8.

=> s hiv-8/ab,bi

1308 HHV/BI  
749460 8/BI  
5408175 AB/FA  
396 HHV-8/AB  
(HHV(V)8)BI (L) AB/FA)  
1308 HHV/BI  
749460 8/BI  
458 HHV-8/BI  
(HHV(V)8)BI  
L3 458 HHV-8/AB,BI

=> s ll and interleukin#/ab,bi

90769 INTERLEUKIN#/BI  
5408175 AB/FA  
59812 INTERLEUKIN#/AB  
(INTERLEUKIN#/BI (L) AB/FA)  
90769 INTERLEUKIN#/BI  
L4 38 L3 AND INTERLEUKIN#/AB,BI

=> s ll and il-6/ab,bi

87055 IL/BI  
1135699 6/BI

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5408175 AB/F A

14922 IL-6/AB

(IL/6/6)BI (L) AB/F A)

87055 IL/BI

1135699 6/BI

15169 IL-6/BI

(IL/6/6)BI)

L5 16 L4 AND IL-6/AB, BI

=> d l - bib ab

YOU HAVE REQUESTED DATA FROM 16 ANSWERS -  
CONTINUE? Y(N)Y

L5 ANSWER 1 OF 16 MEDLINE

AN 2000405417 MEDLINE

DN 20321451

TI Human herpesvirus 8-encoded \*\*\*interleukin\*\*\* -6 homologue  
(viral

\*\*\*TL\*\*\* - \*\*\*6\*\*\* ) induces endogenous human \*\*\*TL\*\*\*  
- \*\*\*6\*\*\*

secretion.

AU Mori Y, Nishimoto N, Ohno M, Inagi R, Dhapakson P, Amonu

K, Yoshizaki K,

Yamanishi K

CS Department of Microbiology, Osaka University Medical School,

Osaka

University Medical School, Osaka University, Osaka, Japan.

ymori@micro.med.osaka-u.ac.jp

SO JOURNAL OF MEDICAL VIROLOGY, (2000 Jul) 61 (3)

332-5.

Journal code: J9N. ISSN: 0146-6615.

CY United States

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 20001004

EW 20001004

AB We found that human herpesvirus 8-encoded \*\*\*TL\*\*\* -

\*\*\*6\*\*\* (vIL-6)

induced endogenous human \*\*\*TL\*\*\* - \*\*\*6\*\*\* (hull-6)

secretion from

various cell lines (MT-4, THP-1, U937, Raji, and CESS) including

patients

with multicentric Castlemans disease. Especially, in MT-4 cells,

hull-6

was enhanced with vIL-6 by 30-fold compared with that of control.

In

addition, reverse transcriptase-polymerase chain reaction confirmed

that

vIL-6 induced hull-6 expression in MT-4 cells. Our novel finding

of the

hull-6 induction by vIL-6 indicates that vIL-6 may play a

significant role

in the pathogenesis of \*\*\*HHV\*\*\* - \*\*\*6\*\*\* associated

diseases.

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L5 ANSWER 2 OF 16 MEDLINE

AN 2000363450 MEDLINE

DN 20363450

TI Presence of human herpesvirus 8 DNA sequences and

overexpression of human

\*\*\*TL\*\*\* - \*\*\*6\*\*\* and cyclin D1 in inflammatory

myofibroblastic tumor

(inflammatory pseudotumor).

AU Gomez-Roman J J, Osejo-Virgala G, Sanchez-Velasco P, Nieto

E H.

Leyva-Cobian F, Val-Bernal J F

CS Departamento de Anatomia Patologica, Hospital Universitario

Marques de

Valdecilla, Instituto Nacional de la Salud, Facultad de Medicina,

Universidad de Cantabria, Santander, Spain.

SO LABORATORY INVESTIGATION, (2000 Jul) 80 (7) 1121-6.

Journal code: KZ4. ISSN: 0023-6837.

CY United States

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals; Cancer Journals

EM 20001001

EW 20001001

AB Inflammatory myofibroblastic tumor (IMT) is composed of

plasma cells, and lymphocytes. Cytokines are possibly involved in

its

pathogenesis. Human herpesvirus-8 ( \*\*\*HHV\*\*\* - \*\*\*6\*\*\* )

encodes cell

cycle regulatory and signaling proteins. A combination of nested

PCR with

several negative controls and Southern blot methods showed the

presence of

\*\*\*HHV\*\*\* - \*\*\*6\*\*\* DNA in seven cases of IMT.

Additionally, strong

expression was demonstrated by in situ hybridization in many

tumoral

nuclei. Most of the myofibroblasts in all of the cases were

immunoreactive

for human \*\*\*TL\*\*\* - \*\*\*6\*\*\* and cyclin D1. These

cytokines probably

have a paracrine action and may sustain myofibroblastic growth.

\*\*\*HHV\*\*\* - \*\*\*6\*\*\* could play an essential role in

triggering IMT

development by a local reactivation of viral lytic replication. The

relationship between \*\*\*HHV\*\*\* - \*\*\*6\*\*\* and

immunosuppression

status as the only associated cause for tumorigenesis should be

revised.

L5 ANSWER 3 OF 16 MEDLINE

AN 2000122659 MEDLINE

DN 20122659

TI Castlemans disease.

AU Palestro G, Turini F, Pegano M, Chiusa L

CS Department of Biomedical Science and Human Oncology,

University of

Torino, Torino, I-10126, Italy. palestro@molinetto.unito.it

SO Adv Clin Path. (1999 Jan-Apr) 3 (1-2) 11-22. Ref: 60

Journal code: DDO. ISSN: 1125-5552.

CY Italy

DT Journal; Article; (JOURNAL ARTICLE)

General Review; (REVIEW)

(REVIEW, TUTORIAL)

LA English

FS Priority Journals

EM 2000004

EW 200000403

AB Castlemans disease (CD) is a rare atypical lymphoproliferative

disorder

whose morphology, soon after the original presentation of

Castlemans et

al, has been definitely subdivided in a hyaline vascular (HV) and

plasma

cell (PC) histopathological pattern, with intermediate variants. The

former occurs much more frequently than the latter and is usually

localized to the mediastinum or pulmonary hilum. The latter

involves lymph

nodes separately or in aggregations and often displays

multicentricity

with systemic symptoms including autoimmune phenomena and

aggressive

course. Infections are the most frequent causes of patient demise in

these

cases, followed by malignancies such as Kaposi's sarcoma,

malignant

lymphoma or epithelial neoplasia. Increase of follicular dendritic

reticulum cells (FDR), often dysplastic, in the germinal center

(GC) and

marginal zone (MZ), broad MZ expansion with prominence of

immunophenotypically aberrant B cells (Ki B3-negative,

CD5-positive),

possible predominance of paracortical plasma cells often with

clusters of

clonal I-light chain restricted plasma cells, increase of paracortical

plasmacytoid monocytes, represent common hallmarks of CD.

However, small

hyalinized and hypervascular GCs with hypervascular interfollicular

stroma

and sinus effacement are common features of the HV variant,

whereas

hyperplastic GCs with plasma cell aggregates in lymph node

paracortex and

partially spaced sinuses are characteristic features of the PC variant.

The frequent concomitance of the HV and PC types at separate

sites,

together with transient morphological patterns from one type to the

other

and from the localized to multicentric form during the course of the

disease, along with B and T cell impaired functions, with frequent

development of autoantibodies, have suggested that CD is a single

disorder

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related to immune dysregulation. A key event in the pathogenesis of CD has been recently suggested to be an abnormal production of a B cell growth factor, such as \*\*\*TL\*\*\* - \*\*\*6\*\*\*, leading to lymphoproliferation and plasma cell differentiation and being involved in the oncogenesis of plasmacytoma. In this event, Kaposi's sarcoma associated virus (\*\*\*HHV\*\*\* - \*\*\*8\*\*\*), which has been found in many cases of CD, especially in the multicentric form, could play a crucial role both in producing \*\*\*TL\*\*\* - \*\*\*6\*\*\* and releasing angiogenic factors. A possible differentiation block may lead to the development of a malignant lymphoma. Kaposi's sarcoma or other malignant neoplasias can be supposed to be consequences of the immunodeficiency typical of CD.

L5 ANSWER 4 OF 16 MEDLINE  
AN 2000053987 MEDLINE  
DN 20053987  
TI Human \*\*\*interleukin\*\*\* -6 is in vivo an autocrine growth factor for human herpesvirus-8-infected malignant B lymphocytes.  
AU Fousat A, Wijdenes J, Bouchet L, Gaidano G, Neipel F, Balabanian K, Galanard P, Couderc J, Emilie D  
CS INSERM U. 131, Institut Paris-Stud sur les Cytokines, 32, rue des Carmes, 92140 Clamart, France  
SO EUROPEAN CYTOKINE NETWORK, (1999 Dec) 10 (4) 501-8.  
Journal code: A56. ISSN: 1148-5493.  
CY France  
DT Journal, Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 200004  
EW 20000402  
AB Human \*\*\*interleukin\*\*\* -6 (hIL-6) acts as a growth factor in several human B lymphoid cancers. As human herpesvirus-8 (\*\*\*HHV\*\*\* - \*\*\*8\*\*\*) encodes for a viral \*\*\*TL\*\*\* - \*\*\*6\*\*\* (vIL-6), the viral cytokine may be responsible for several manifestations of \*\*\*HHV\*\*\* - \*\*\*8\*\*\*-related disorders. Using an anti-hIL-6 mAb (B-E8) which does not recognize vIL-6, we investigated the involvement of the human cytokine in the proliferation of \*\*\*HHV\*\*\* - \*\*\*8\*\*\*-positive primary effusion lymphoma (PEL) cells. In vitro, S/S PEL cell lines produced hIL-6 (4 to 1,200 pg/ml). The EBV- \*\*\*HHV\*\*\* - \*\*\*8\*\*\* + cell line

(BCBL-1) was adapted to grow in SCID mice. hIL-6 was detected in the serum of mice with grafts, as well as human soluble CD138 (sCD138) and human IL-10 (hIL-10). The serum level of these mediators increased with tumor progression. The effect of treatment with the B-E8 mAb on the tumor progression and survival was evaluated. This treatment significantly slowed down the tumor development: on day 54, there were more mice with low levels of sCD138 and hIL-10 in the treated group than in controls (p = 0.03 and 0.02, respectively); treatment also delayed death (median date of death was day 65 for control mice and day 84 for anti-hIL-6 mAb-treated mice; p < 0.02). Thus, hIL-6 is expressed in addition to vIL-6 in \*\*\*HHV\*\*\* - \*\*\*8\*\*\*-positive malignant B lymphocytes, and the viral cytokine does not substitute for human \*\*\*TL\*\*\* - \*\*\*6\*\*\* in promoting tumor progression.

L5 ANSWER 5 OF 16 MEDLINE  
AN 1999445405 MEDLINE  
DN 99445405  
TI Expression of cell-homologous genes of human herpesvirus-8 in human immunodeficiency virus-negative lymphoproliferative diseases.  
AU Luppi M, Barozzi P, Maiorana A, Trovato R, Marasca R, Morselli M, Caggisi K, Torelli G  
CS Department of Medical Sciences, Section of Hematology, Modena, Italy.  
SO BLOOD, (1999 Oct 15) 94 (8) 2931-3.  
Journal code: A8G. ISSN: 0006-4971.  
CY United States  
DT Journal, Article; (JOURNAL ARTICLE)  
LA English  
FS Abstracted Index Medicus Journals; Priority Journals; Cancer Journals  
EM 200001  
EW 20000104  
AB Human herpesvirus-8 (\*\*\*HHV\*\*\* - \*\*\*8\*\*\*) genome encodes for genes homologous to human cellular genes such as \*\*\*interleukin\*\*\* -6 (\*\*\*TL\*\*\* - \*\*\*6\*\*\*), Cyclin-D, BCL-2, and IL-8 receptor (G-protein-coupled receptor [GCR]). We used reverse transcriptase-polymerase chain reaction to study the expression of these viral genes in lymphoproliferative disorders associated with \*\*\*HHV\*\*\* - \*\*\*8\*\*\* infection. None of these genes was expressed in 1 case of benign,

localized Castleman's disease (CD), and only viral \*\*\*TL\*\*\* - \*\*\*6\*\*\* and viral Cyclin-D were transcribed in 2 cases of benign lymphadenopathies with giant germinal center hyperplasia and increased vascularity. In contrast, all 4 genes were transcribed in 1 case of multicentric CD of plasma cell type with aggressive clinical course and in 1 primary effusion lymphoma cell line. Our study provides the evidence that various \*\*\*HHV\*\*\* - \*\*\*8\*\*\* genes, homologous to cellular genes involved in control of proliferation and apoptosis, may be differently expressed in different lymphoid disorders in vivo.

L5 ANSWER 6 OF 16 MEDLINE  
AN 1999441800 MEDLINE  
DN 99441800  
TI Human herpesvirus-8-encoded \*\*\*interleukin\*\*\* 6 activates HIV-1 in the monocytic cell line.  
AU Gage J R, Breen E C, Baliverni A, Maggarity L, Kishimoto T, Miles S, Martinez-Maza O  
CS Department of Obstetrics and Gynecology, Osaka University, Japan.  
NC CA57152 (NCI)  
CA73475 (NCI)  
F32CA80610 (NCI)  
+  
SO AIDS, (1999 Oct 1) 13 (14) 1851-5.  
Journal code: AID. ISSN: 0269-9370.  
CY ENGLAND: United Kingdom  
DT Journal, Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 200002  
EW 20000204  
AB OBJECTIVE: Human herpesvirus 8 (\*\*\*HHV\*\*\* - \*\*\*8\*\*\*) encodes a viral \*\*\*interleukin\*\*\* 6 (vIL-6) which is structurally and functionally similar to human \*\*\*interleukin\*\*\* 6 (hIL-6). Since hIL-6 has been shown to upregulate the expression of HIV-1, the objectives of this study were to examine the ability of vIL-6 to upregulate HIV-1, and to determine the interactions of this cytokine (viral cytokine) with the components of the \*\*\*interleukin\*\*\* 6 (\*\*\*TL\*\*\* - \*\*\*6\*\*\*) receptor complex.  
DESIGN AND METHODS: Recombinant \*\*\*HHV\*\*\* - \*\*\*8\*\*\* vIL-6 (vIL-6) was assayed for bioactivity in the \*\*\*TL\*\*\* - \*\*\*6\*\*\*-dependent cell

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line MH60.BSF2. HIV-1 p24 production by the U1 monocytic and ACH-2 T-cell lines, which are chronically infected with HIV-1, was used to assess the ability of vIL-6 to affect HIV-1 expression, hIL-6 and vIL-6 receptor utilization was determined by quantifying HIV-1 p24 production after neutralization of components of the \*\*\*IL\*\*\*. \*\*\*6\*\*\* receptor complex, CD126IL-6R and CD130gp130, on U1 cells with blocking antibodies. RESULTS: \*\*\*HHV\*\*\*. \*\*\*6\*\*\* vIL-6 was seen to have \*\*\*IL\*\*\*. \*\*\*6\*\*\*-like bioactivity in MH60.BSF2 cells, and readily upregulated HIV-1 p24 production in U1 monocytic cells, but not in ACH-2 T cells. The vIL-6 appeared to utilize the \*\*\*IL\*\*\*. \*\*\*6\*\*\*-specific component of the \*\*\*IL\*\*\*. \*\*\*6\*\*\* signaling complex, CD126IL-6R, in U1 cells. CONCLUSIONS: \*\*\*HHV\*\*\*. \*\*\*6\*\*\* vIL-6 clearly has the potential to upregulate HIV-1 expression in monocytic cells, and therefore may play a role in AIDS pathogenesis in individuals infected with both viruses.

L5 ANSWER 7 OF 16 MEDLINE  
AN 1999412342 MEDLINE  
DN 99412342  
TI Human herpesvirus 8 \*\*\*interleukin\*\*\*-6 (vIL-6) signals through gp130 but has structural and receptor-binding properties distinct from those of human \*\*\*IL\*\*\*. \*\*\*6\*\*\*  
AU Man X, Wang H, Nicholas J  
CS Molecular Virology Laboratories, Department of Oncology, Johns Hopkins University School of Medicine, Baltimore, Maryland 21231, USA.  
NC R53 CA76445 (NCI)  
SO JOURNAL OF VIROLOGY, (1999 Oct) 73 (10) 8268-78.  
Journal code: KCV. ISSN: 0022-538X.  
CY United States  
DT Journal Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals; Cancer Journals  
EM 199912  
EW 19991203  
AB Human herpesvirus 8 ( \*\*\*HHV\*\*\*. \*\*\*6\*\*\* ) has been associated with classical, endemic (African), and AIDS-related Kaposi's sarcoma (KS), body cavity-based primary effusion lymphomas, and multicentric

Castleman's disease (MCD). \*\*\*HHV\*\*\*. \*\*\*6\*\*\* encodes a functional homologue of \*\*\*interleukin\*\*\*-6 ( \*\*\*IL\*\*\*. \*\*\*6\*\*\* ), a cytokine that promotes the growth of KS and myeloma cells and is found at elevated levels in MCD lesions and patient sera. We have previously reported that the viral \*\*\*IL\*\*\*. \*\*\*6\*\*\* (vIL-6) gene product can support the growth of the \*\*\*IL\*\*\*. \*\*\*6\*\*\*-dependent murine hybridoma cell line, B9, and that the gp80 ( \*\*\*IL\*\*\*. \*\*\*6\*\*\* receptor [IL-6R]) component of the \*\*\*IL\*\*\*. \*\*\*6\*\*\* receptor-signal transducer (gp180) complex plays a role in mediating this activity. However, it has been shown by others that vIL-6 can function in human cells independently of IL-6R. Here we have extended our functional studies of vIL-6 by identifying transcription factors and pathways used in human Hep3B cells, investigating the utilization of gp130 and IL-6R by vIL-6, and undertaking mutational analyses of vIL-6 and gp130. The data presented here establish that vIL-6, in common with its endogenous counterparts, can mediate signal transduction through gp130 and activate multiple transcription factors, map residues within the vIL-6 protein that are and are not important for vIL-6 signalling, and identify a gp130 mutant that is nonfunctional with respect to vIL-6 signalling in the absence of IL-6R but that retains the ability to mediate vIL-6 and human \*\*\*IL\*\*\*. \*\*\*6\*\*\* (hIL-6) signal transduction when IL-6R is coexpressed. The data presented demonstrate functional and mechanistic similarities between vIL-6 and endogenous \*\*\*IL\*\*\*. \*\*\*6\*\*\* proteins but also highlight differences in the structural and receptor-binding properties of vIL-6 relative to its human counterpart.

L5 ANSWER 8 OF 16 MEDLINE  
AN 1999367590 MEDLINE  
DN 99367590  
TI Heterogeneity of viral \*\*\*IL\*\*\*. \*\*\*6\*\*\* expression in \*\*\*HHV\*\*\*  
\*\*\*6\*\*\*-associated diseases.  
AU Cannon JS, Nicholas J, Orenstein J M, Mann R B, Murray P G, Browning P J,  
DiGiuseppe J A, Cesarman E, Hayward G S, Ambinder R F  
CS Department of Pharmacology, Johns Hopkins University School

of Medicine, Baltimore, Maryland USA.  
NC U01 CA70062 (NCI)  
PO30 CA06973 (NCI)  
PO30 CA68485 (NCI)  
SO JOURNAL OF INFECTIOUS DISEASES, (1999 Sep) 180 (3) 824-8.  
Journal code: IH3. ISSN: 0022-1899.  
CY United States  
DT Journal Article; (JOURNAL ARTICLE)  
LA English  
FS Abridged Index Medicus Journals; Priority Journals  
EM 19991105  
EW 19991105  
AB In order to characterize the expression of the viral \*\*\*interleukin\*\*\*-6 (vIL-6) homologue in various human herpesvirus 8 ( \*\*\*HHV\*\*\*. \*\*\*6\*\*\* )-associated diseases, in situ hybridization and immunohistochemistry were applied to formalin-fixed specimens. These assays showed consistent expression of vIL-6 in primary effusion lymphomas and in a case of human immunodeficiency virus (HIV)-associated lymphadenopathy with a Castleman's disease-like appearance. In contrast, Kaposi's sarcoma specimens showed marked differences among specimens. In a consecutive series of specimens from the Johns Hopkins archives, vIL-6 expression was demonstrated in one of 13 cases. However, among 7 specimens selected from the AIDS Malignancy Bank because of their high levels of the T.1.1 lytic transcript and viron production, vIL-6 expression was consistently demonstrated in infiltrating mononuclear cells and occasional spindle-shaped cells. Thus vIL-6 expression in clinical specimens correlates with other measures of the lytic viral cycle. Both assays generally give congruent results and are consistent with the possibility that vIL-6 expression plays a role in the pathogenesis of a variety of \*\*\*HHV\*\*\*. \*\*\*6\*\*\*-associated diseases.

L5 ANSWER 9 OF 16 MEDLINE  
AN 1999292923 MEDLINE  
DN 99292923  
TI A tlesu maeque thadimovirus related to Kaposi's sarcoma-associated herpesvirus/human herpesvirus 8 encodes a functional homologue of \*\*\*interleukin\*\*\*-6.  
AU Kaleeba J A, Bergum E P, Wong S W  
CS Division of Pathobiology and Immunology, Oregon Regional Primate Research Center, Beaverton, Oregon 97006, USA  
NC CA75922 (NCI)

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\*\*\*IL\*\*\* . \*\*\*6\*\*\* , IL-10, IL-11, LIF, MIP-1alpha, or OSM. Inhibition with IL-6R and \*\*\*IL\*\*\* . \*\*\*6\*\*\* inhibited cell growth. Anti-IL-6 neutralizing antibodies had no effect on PEL cell line proliferation, conversely, whereas anti-IL-6R alone inhibited only weakly, anti-gp130 and anti-gp130 plus anti-IL-6R showed strong inhibitory effects (>20% inhibition in 5/9 lines and >60% inhibition in 3/9 lines). In summary, PEL cell lines produce high amounts of cytokines ( \*\*\*IL\*\*\* . \*\*\*6\*\*\* IL-10, OSM), proliferation could be inhibited by blocking the receptors of the \*\*\*IL\*\*\* . \*\*\*6\*\*\* signaling pathway.

L5 ANSWER 12 OF 16 MEDLINE  
AN 1998374965 MEDLINE  
DN 98374965  
TI Human herpesvirus type 8 and Kaposi's sarcoma  
AU Weiss R A, Whitty D, Talbot S, Kellam P, Boshoff C  
CS Institute of Cancer Research, Chester Beatty Laboratories, London, U.K.  
SO JOURNAL OF THE NATIONAL CANCER INSTITUTE.  
MONOGRAPHS. (1998) (23) 51-4.  
Ref: 32  
Journal code: ATR. ISSN: 1052-6773.  
CY United States  
DT Journal Article. (JOURNAL ARTICLE)  
General Review. (REVIEW)  
(REVIEW, TUTORIAL)  
LA English  
FS Priority Journals  
EM 19981102  
AB Kaposi's sarcoma-associated herpesvirus or human herpesvirus type 8 ( \*\*\*HHV\*\*\* . \*\*\*8\*\*\* ) is present in all forms of Kaposi's sarcoma (KS) as well as in primary effusion lymphomas and some cases of Castleman's disease. In KS tissues, \*\*\*HHV\*\*\* . \*\*\*8\*\*\* is present in endothelial and spindle cells. Current serologic tests suggest that \*\*\*HHV\*\*\* . \*\*\*8\*\*\* is predominantly found in those at risk of KS and is not as widespread as most other human herpesviruses. \*\*\*HHV\*\*\* . \*\*\*8\*\*\* encodes various proteins that may play a role in promotion of cellular growth, including cyclin- and G-coupled protein receptor homologues, and anti-apoptotic proteins, including Bcl-2. \*\*\*IL\*\*\* . \*\*\*6\*\*\* (i.e., \*\*\*interleukin\*\*\* 6), and FLIP (i.e., FLICE inhibitory protein) homologues. In addition, \*\*\*HHV\*\*\* . \*\*\*8\*\*\* encodes two macrophage inflammatory-like proteins with anti-human

immunodeficiency virus and angiogenic potential.

L5 ANSWER 13 OF 16 MEDLINE  
AN 1998158619 MEDLINE  
DN 98158619  
TI Human herpesvirus type 8 \*\*\*interleukin\*\*\* -6 homologue is functionally active on human myeloma cells [see comments]  
CM Comment in: Blood 1998 Sep 15;92(6):2186-8  
AU Burger R, Neipel F, Fleckenstein B, Savino R, Ciliberto G, Kalden J R, Gramatz M  
CS Division of Hematology/Oncology, Department of Medicine III, the Institute for Clinical and Molecular Virology, University of Erlangen-Nuremberg, Erlangen, Germany,  
SO BLOOD. (1998 Mar 15) 91 (6) 1838-63.  
Journal code: A8G. ISSN: 0006-4971.  
CY United States  
DT Journal Article. (JOURNAL ARTICLE)  
LA English  
FS Abridged Index Medicus Journals; Priority Journals; Cancer Journals  
EM 199806  
AB Seroprevalence and polymerase chain reaction studies have strongly suggested that human herpesvirus type 8 ( \*\*\*HHV\*\*\* . \*\*\*8\*\*\* ) is associated with Kaposi's sarcoma, Castleman's disease, and body cavity-based lymphoma. The genome of \*\*\*HHV\*\*\* . \*\*\*8\*\*\* harbors a viral analogue of the \*\*\*interleukin\*\*\* -6 ( \*\*\*IL\*\*\* . \*\*\*6\*\*\* ) gene. The amino acid sequence of the viral \*\*\*IL\*\*\* . \*\*\*6\*\*\* (vIL-6) protein is 24.7% identical to human \*\*\*IL\*\*\* . \*\*\*6\*\*\* (hIL-6). \*\*\*IL\*\*\* . \*\*\*6\*\*\* as a B-cell growth and differentiation factor is known to play an essential role in the pathophysiology of B-cell tumors. Thus, it seems possible that virus-encoded \*\*\*IL\*\*\* . \*\*\*6\*\*\* contributes to malignant growth of \*\*\*HHV\*\*\* . \*\*\*8\*\*\* -positive B-cell lymphatic tumors. We have tested a preparation of \*\*\*HHV\*\*\* . \*\*\*8\*\*\* -derived \*\*\*IL\*\*\* . \*\*\*6\*\*\* for the ability to promote the proliferation of the human myeloma cell line INA-6, which is strictly dependent on exogenous \*\*\*IL\*\*\* . \*\*\*6\*\*\* for growth and survival. Viral \*\*\*IL\*\*\* . \*\*\*6\*\*\* significantly induced DNA synthesis of INA-6 cells, but required much more protein on a weight basis

when compared with hIL-6 for maximal proliferation. The proliferative effect of vIL-6 was almost completely inhibited by a combination of anti-\*\*\*IL\*\*\* . \*\*\*6\*\*\* receptor (IL-6R) and anti-gp130 antibodies or IL-6R superantagonist Sant7 and anti-gp130 antibodies. This report demonstrates that vIL-6 has proliferative activity on human cells and that the IL-6R and gp130 are involved in vIL-6 signaling in the myeloma cell line INA-6.

L5 ANSWER 14 OF 16 MEDLINE  
AN 97208913 MEDLINE  
DN 97208913  
TI Kaposi's sarcoma-associated human herpesvirus-8 encodes homologues of macrophage inflammatory protein-1 and \*\*\*interleukin\*\*\* -6.  
AU Nicholas J, Rutvick V R, Burns W H, Sandford G, Wan X, Cutro D, Hendrickson S B, Guo H G, Hayward G S, Reitz M S  
CS Department of Oncology, Johns Hopkins University School of Medicine, Baltimore, Maryland 21205, USA.  
SO NATURE MEDICINE. (1997 Mar) 3 (3) 287-92.  
Journal code: CG5. ISSN: 1078-8956.  
CY United States  
DT Journal Article. (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
OS GENBANK-U67774; GENBANK-U67775; GENBANK-U74585  
EM 199706  
AB Human herpesvirus-8 ( \*\*\*HHV\*\*\* . \*\*\*8\*\*\* ) has been detected in Kaposi's sarcoma (KS) lesions of all types (AIDS-related, classical and endemic), in body-cavity-based B-cell lymphomas (BCBLs) and in lesions of multicentric Castleman's disease (MCD). We have identified a major gamma-herpesvirus-divergent locus (DL-B) in \*\*\*HHV\*\*\* . \*\*\*8\*\*\* DNA encoding several \*\*\*HHV\*\*\* . \*\*\*8\*\*\* unique open reading frames (ORFs), including a homologue of \*\*\*interleukin\*\*\* -6 ( \*\*\*IL\*\*\* . \*\*\*6\*\*\* ) and two homologues of macrophage inflammatory protein MIP-1. We show that the \*\*\*HHV\*\*\* . \*\*\*8\*\*\* -encoded \*\*\*IL\*\*\* . \*\*\*6\*\*\* homologue (vIL-6) shares functional properties with endogenous \*\*\*IL\*\*\* . \*\*\*6\*\*\* proteins and that both vIL-6 and vMIP-1 transcripts are present at high levels following butyrate induction of an

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28829 HERPESVIRUS/BI  
52956 TYPE/BI  
74940 8/BI  
57 HUMAN HERPESVIRUS TYPE 8/BI  
(HUMAN(V)HERPESVIRUS(W)TYPE(W)8/BI)  
L6 57 HUMAN HERPESVIRUS TYPE 8/AB,BI  
=> s16 and interleukin##/ab,bi  
90769 INTERLEUKIN##/BI  
5408175 AB/FA  
59812 INTERLEUKIN##/AB  
(INTERLEUKIN##/BI (L) AB/FA)  
L7 90769 INTERLEUKIN##/BI  
416 AND INTERLEUKIN##/AB,BI  
=> d1 - bib ab  
YOU HAVE REQUESTED DATA FROM 4 ANSWERS.  
CONTINUE? Y(N)?  
L7 ANSWER 1 OF 4 MEDLINE  
AN 1999316867 MEDLINE  
DN 99316867  
TI The \*\*\*human\*\*\* \*\*\*herpesvirus\*\*\* - \*\*\*type\*\*\*  
\*\*\*\*g\*\*\* is  
not involved in malignant melanoma.  
AU Deichmann M, Thome M, Beck M, Jacek A, Waldmann V,  
Naher H  
CS Universitäts-Haarklinik Heidelberg, Germany.  
SO BRITISH JOURNAL OF CANCER. (1999 Apr) 80 (1-2) 67-9.  
Journal code: AV4. ISSN: 0007-0920.  
CY SCOTLAND: United Kingdom  
DT Journal Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals; Cancer Journals  
EM 199909  
EW 19990902  
AB Malignant melanomas were supposed to harbour the  
\*\*\*human\*\*\*  
\*\*\*herpesvirus\*\*\* - \*\*\*type\*\*\* \*\*\*\*g\*\*\* (HHV-8)  
genome, as melanoma  
cells were reported to express \*\*\*interleukin\*\*\* -6 and a  
homologue of  
\*\*\*interleukin\*\*\* -6 was found in the HHV-8 genome. We  
therefore  
investigated 33 primary malignant melanomas by polymerase chain  
reaction,  
but could not find this tumorigenic gamma-herpesvirus in any  
tumour.  
L7 ANSWER 2 OF 4 MEDLINE  
AN 1999223975 MEDLINE  
DN 99223975  
TI Involvement of human \*\*\*interleukin\*\*\* -6 in systemic  
manifestations of

\*\*\*human\*\*\* \*\*\*herpesvirus\*\*\* \*\*\*type\*\*\* \*\*\*\*g\*\*\*  
-associated  
multicentric Castlemans disease [letter].  
AU Fousas A, Fior R, Girard T, Boue F, Wijdenes J, Calanard P,  
Emile D  
SO AIDS. (1999 Jan 14) 13 (1) 150-2.  
Journal code: AID. ISSN: 0269-9370.  
CY ENGLAND: United Kingdom  
DT Letter  
LA English  
FS Priority Journals  
EM 199909  
EW 19990901  
L7 ANSWER 3 OF 4 MEDLINE  
AN 1998374965 MEDLINE  
DN 98374965  
TI \*\*\*Human\*\*\* \*\*\*herpesvirus\*\*\* \*\*\*type\*\*\*  
\*\*\*\*g\*\*\* and  
Kaposi's sarcoma.  
AU Weiss R A, Whitty D, Talbot S, Kellam P, Boethoff C  
CS Institute of Cancer Research, Chester Beatty Laboratories,  
London, UK.  
SO JOURNAL OF THE NATIONAL CANCER INSTITUTE.  
MONOGRAPHS. (1998) (23) 51-4.  
Ref: 32  
Journal code: ATR. ISSN: 1052-6773.  
CY United States  
DT Journal Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, TUTORIAL)  
LA English  
FS Priority Journals  
EM 199811  
EW 19981102  
AB Kaposi's sarcoma-associated herpesvirus or \*\*\*human\*\*\*  
\*\*\*herpesvirus\*\*\* \*\*\*type\*\*\* \*\*\*\*g\*\*\* (HHV-8) is  
present in all  
forms of Kaposi's sarcoma (KS) as well as in primary effusion  
lymphomas  
and some cases of Castlemans disease. In KS tissues, HHV-8 is  
present in  
endothelial and spindle cells. Current serologic tests suggest that  
HHV-8  
is predominantly found in those at risk of KS and is not as  
widespread as  
most other human herpesviruses. HHV-8 encodes various proteins  
that may  
play a role in promotion of cellular growth, including cyclin- and  
G-coupled protein receptor homologues, and anti-apoptotic  
proteins,  
including Bcl-2, IL-6 (i.e., \*\*\*interleukin\*\*\* -6), and FLIP (i.e.,  
FLICE inhibitory protein) homologues. In addition, HHV-8  
encodes two  
macrophage inflammatory-like proteins with anti-human  
immunodeficiency  
virus and angiogenic potential.

L7 ANSWER 4 OF 4 MEDLINE  
AN 1998158619 MEDLINE  
DN 98158619  
TI \*\*\*Human\*\*\* \*\*\*herpesvirus\*\*\* \*\*\*type\*\*\*  
\*\*\*\*g\*\*\*  
\*\*\*interleukin\*\*\* -6 homologue is functionally active on human  
myeloma  
cells [see comments].  
CM Comment in: Blood 1998 Sep 15;92(6):2186-8  
AU Burger R, Neipel F, Fleckenstein B, Servino R, Ciliberto G,  
Kalden J R,  
Granatzki M  
CS Division of Hematology/Oncology, Department of Medicine III,  
the Institute  
for Clinical and Molecular Virology, University of  
Erlangen-Nuernberg,  
Erlangen, Germany.  
SO BLOOD. (1998 Mar 15) 91 (6) 1858-63.  
Journal code: A8G. ISSN: 0006-4971.  
CY United States  
DT Journal Article; (JOURNAL ARTICLE)  
LA English  
FS Abridged Index Medicus Journals; Priority Journals; Cancer  
Journals  
EM 199806  
AB Seropidemiology and polymerase chain reaction studies have  
strongly  
suggested that \*\*\*human\*\*\* \*\*\*herpesvirus\*\*\*  
\*\*\*type\*\*\*  
\*\*\*\*g\*\*\* (HHV-8) is associated with Kaposi's sarcoma,  
Castlemans  
disease, and body cavity-based lymphoma. The genome of HHV-8  
harbors a  
viral analogue of the \*\*\*interleukin\*\*\* -6 (IL-6) gene. The  
amino acid  
sequence of the viral IL-6 (vIL-6) protein is 24.7% identical to  
human  
IL-6 (hIL-6). IL-6 as a B-cell growth and differentiation factor is  
known  
to play an essential role in the pathophysiology of B-cell tumors.  
Thus,  
it seems possible that virus-encoded IL-6 contributes to malignant  
growth  
of HHV-8-positive B-cell lymphatic tumors. We have tested a  
preparation of  
HHV-8-derived IL-6 for the ability to promote the proliferation of  
the  
human myeloma cell line INA-6, which is strictly dependent on  
exogenous  
IL-6 for growth and survival. Viral IL-6 significantly induced DNA  
synthesis of INA-6 cells, but required much more protein on a  
weight basis  
when compared with hIL-6 for maximal proliferation. The  
proliferative  
effect of vIL-6 was almost completely inhibited by a combination  
of

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anti-IL-6 receptor (IL-6R) and anti-gp130 antibodies or IL-6R superantagonist Sant7 and anti-gp130 antibodies. This report demonstrates that VIL-6 has proliferative activity on human cells and that the IL-6R and gp130 are involved in VIL-6 signaling in the myeloma cell line INA-6.

=> d his

(FILE HOME ENTERED AT 16:30:20 ON 15 SEP 2000)

FILE MEDLINE ENTERED AT 16:30:25 ON 15 SEP 2000  
L1 14 S HERPES VIRUS TYPE 8/AB.BI  
L2 1 S L1 AND INTERLEUKIN/AB.BI  
L3 458 S HHV-8/AB.BI  
L4 38 S L3 AND INTERLEUKIN/AB.BI  
L5 16 S L4 AND IL-6/AB.BI  
L6 57 S HUMAN HERPESVIRUS TYPE 8/AB.BI  
L7 4 S L6 AND INTERLEUKIN/AB.BI

=> file medicine embase biosis impadoc caplus

COST IN U.S. DOLLARS SINCE FILE TOTAL  
ENTRY SESSION  
FULL ESTIMATED COST 5.74 5.89

FILE MEDLINE ENTERED AT 16:33:56 ON 15 SEP 2000

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=> e fleckenstein bernhard/au

E1 9 FLECKENSTEIN BERNHARD/AU  
E2 4 FLECKENSTEIN BERNHARD PROF/AU  
E3 235 -> FLECKENSTEIN BERNHARD/AU  
E4 1 FLECKENSTEIN BERNHARD DR/AU  
E5 1 FLECKENSTEIN BERNHARD DR PROF/AU  
E6 3 FLECKENSTEIN BERNHARD PROF/AU  
E7 23 FLECKENSTEIN BERNHARD PROF DR/AU  
E8 5 FLECKENSTEIN BERNHARD W/AU  
E9 24 FLECKENSTEIN BURKHARD/AU

E10 8 FLECKENSTEIN BURKHARD T/AU  
E11 10 FLECKENSTEIN CHRISTOPH/AU  
E12 3 FLECKENSTEIN D/AU

=> s e1-e10

L8 312 (\*FLECKENSTEIN BERNHARD\*/AU OR  
\*FLECKENSTEIN BERNHARD PROF\*/AU OR  
\*FLECKENSTEIN BERNHARD\*/AU OR  
\*FLECKENSTEIN BERNHARD DR\*/AU OR  
\*FLECKENSTEIN BERNHARD DR PROF\*/AU OR  
\*FLECKENSTEIN BERNHARD  
PROF\*/AU OR \*FLECKENSTEIN BERNHARD PROF  
DR\*/AU OR \*FLECKENSTEIN  
BERNHARD W\*/AU OR \*FLECKENSTEIN  
BURKHARD\*/AU OR \*FLECKENSTEIN  
BURKHARD T\*/AU)

=> e albrecht jens-christian/au

E1 30 ALBRECHT JENS CHRISTIAN/AU  
E2 2 ALBRECHT JENS DIPL ING/AU  
E3 0 -> ALBRECHT JENS-CHRISTIAN/AU  
E4 1 ALBRECHT JERRY L/AU  
E5 6 ALBRECHT JERRY V/AU  
E6 1 ALBRECHT JIM/AU  
E7 73 ALBRECHT JIRU/AU  
E8 1 ALBRECHT JIRI DIPL ING/AU  
E9 41 ALBRECHT JIRI ING CSC/AU  
E10 55 ALBRECHT JOACHIM/AU  
E11 23 ALBRECHT JOACHIM DR/AU  
E12 2 ALBRECHT JOACHIM DR ING/AU

=> e albrecht jens-christian/au

E1 18 ALBRECHT JENS/AU  
E2 3 ALBRECHT JENS C/AU  
E3 30 -> ALBRECHT JENS CHRISTIAN/AU  
E4 2 ALBRECHT JENS DIPL ING/AU  
E5 1 ALBRECHT JERRY L/AU  
E6 6 ALBRECHT JERRY V/AU  
E7 1 ALBRECHT JIM/AU  
E8 73 ALBRECHT JIRU/AU  
E9 1 ALBRECHT JIRI DIPL ING/AU  
E10 41 ALBRECHT JIRI ING CSC/AU  
E11 55 ALBRECHT JOACHIM/AU  
E12 23 ALBRECHT JOACHIM DR/AU

=> s e1-e3

L9 51 (\*ALBRECHT JENS\*/AU OR \*ALBRECHT JENS  
C\*/AU OR \*ALBRECHT JENS  
CHRISTIAN\*/AU)

=> e neipel frank/au

E1 1 NEIPALU V/AU  
E2 82 NEIPEL F/AU  
E3 62 -> NEIPEL FRANK/AU  
E4 1 NEIPEL FRANK DR/AU  
E5 1 NEIPEL M P/AU  
E6 19 NEIPELT K V/AU  
E7 4 NEIPELT M/AU  
E8 2 NEIPELT M P/AU  
E9 13 NEIPELT MARSHALL P/AU  
E10 1 NEIPELT MARSHALL PALMER/AU  
E11 1 NEIPLING J E/AU  
E12 3 NEIPLING L/AU

=> s e2-e4

L10 145 (\*NEIPEL F\*/AU OR \*NEIPEL FRANK\*/AU OR  
\*NEIPEL FRANK DR\*/AU)

=> e friedman kien alvin/au

E1 79 FRIEDMAN KIEN A/AU  
E2 338 FRIEDMAN KIEN A E/AU  
E3 15 -> FRIEDMAN KIEN AL VIN/AU  
E4 64 FRIEDMAN KIEN AL VIN E/AU  
E5 1 FRIEDMAN KIEN U/AU  
E6 1 FRIEDMAN KIEN A/AU  
E7 6 FRIEDMAN KIM J/AU  
E8 1 FRIEDMAN KIEN A E/AU  
E9 1 FRIEDMAN KIEN A/AU  
E10 2 FRIEDMAN KURT E/AU  
E11 933 FRIEDMAN L/AU  
E12 82 FRIEDMAN L A/AU

=> s e1-e4

L11 496 (\*FRIEDMAN KIEN A\*/AU OR \*FRIEDMAN KIEN A  
E\*/AU OR \*FRIEDMAN  
KIEN AL VIN\*/AU OR \*FRIEDMAN KIEN AL VIN  
E\*/AU)

=> e huang yao qi/au

E1 2 HUANG YAO QI/AU  
E2 3 HUANG YAO Q/AU  
E3 42 -> HUANG YAO QI/AU  
E4 1 HUANG YAO QIAN/AU  
E5 2 HUANG YAO SHEN/AU  
E6 4 HUANG YAO T/AU  
E7 9 HUANG YAO TE/AU  
E8 4 HUANG YAO TSENG/AU  
E9 6 HUANG YAO TSUNG/AU  
E10 4 HUANG YAO WEI/AU  
E11 23 HUANG YAO WEN/AU  
E12 6 HUANG YAO XIONG/AU

=> s e1-e4

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L12 48 (HUANG YAO QI/AU OR HUANG YAO QI/AU OR  
HUANG YAO QI/AU OR  
HUANG YAO QUAN/AU)

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(FILE HOME ENTERED AT 16:30:20 ON 15 SEP 2000)

FILE MEDLINE ENTERED AT 16:30:25 ON 15 SEP 2000

L1 14 S HERPES VIRUS TYPE 8/AB, BI  
L2 1 S L1 AND INTERLEUKIN/AB, BI  
L3 458 S HHV-8/AB, BI  
L4 38 S L3 AND INTERLEUKIN/AB, BI  
L5 16 S L4 AND IL-6/AB, BI  
L6 57 S HUMAN HERPESVIRUS TYPE 8/AB, BI  
L7 4 S L6 AND INTERLEUKIN/AB, BI

FILE MEDLINE, EMBASE, BIOSIS, INPADOC, CAPLUS  
ENTERED AT 16:33:56 ON 15  
SEP 2000

E FLECKENSTEIN BERNHARD/AU  
L8 312 S E1-E10  
E ALBRECHT JENS-CHRISTIAN/AU  
E ALBRECHT JENS CHRISTIAN/AU  
L9 51 S E1-E3  
E NEIPEL FRANK/AU  
L10 145 S E2-E4  
E FRIEDMAN KIEN ALVIN/AU  
L11 456 S E1-E4  
E HUANG YAO QI/AU  
L12 48 S E1-E4

=> s 18 or 19 or 11 or 112

L13 937 L8 OR L9 OR L10 OR L11 OR L12

=> s 113 and (11 or 13 or 16)

'AB IS NOT A VALID FIELD CODE  
L14 82 L13 AND (L1 OR L3 OR L6)

=> s 114 and interleukin/ab, bi

'AB IS NOT A VALID FIELD CODE  
L15 20 L14 AND INTERLEUKIN/AB, BI

=> dup rem 115

PROCESSING COMPLETED FOR L15

L16 10 DUP REM L15 (10 DUPLICATES REMOVED)

=> d 1 - bib ab

YOU HAVE REQUESTED DATA FROM 10 ANSWERS.  
CONTINUE? Y(N)?

L16 ANSWER 1 OF 10 MEDLINE DUPLICATE

AN 2000053987 MEDLINE  
DN 20053987

TI Human \*\*\*interleukin\*\*\*-6 is in vivo an autocrine growth  
factor for

human herpesvirus-8-infected malignant B lymphocytes.

AU Foussea A, Wijdenes J, Bouchet L, Gaidano G, \*\*\*Neipel  
P\*\*\*

Balaban K, Galanard P, Couderc J, Emilie D

CS INSERM U. 131, Institut Paris-Sud sur les Cytokines, 32, rue des  
Carnets,

92140 Clamart, France.

SO EUROPEAN CYTOKINE NETWORK. (1999 Dec) 10 (4)  
501-8.

Journal code: A56 ISSN: 1148-5493.

CY France

DT Journal Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 200004

EW 20000402

AB Human \*\*\*interleukin\*\*\*-6 (IL-6) acts as a growth factor in  
several

human B lymphoid cancers. As human herpesvirus-8 (

\*\*\*HHV\*\*\*-8

) encodes for a viral IL-6 (vIL-6), the viral cytokine may be

responsible

for several manifestations of \*\*\*HHV\*\*\*-8-related

disorders.

Using an anti-IL-6 mAb (B-E8) which does not recognize vIL-6,

we

investigated the involvement of the human cytokine in the

proliferation of

\*\*\*HHV\*\*\*-8-positive primary effusion lymphoma

(PEL) cells. In

vitro, 5/5 PEL cell lines produced hIL-6 (4 to 1,200 pg/ml). The

EBV-

\*\*\*HHV\*\*\*-8 + cell line (BCBL-1) was adapted to

grow in SCID

mice. hIL-6 was detected in the serum of mice with grafts, as well

as

human soluble CD138 (sCD138) and human IL-10 (hIL-10). The

serum level of

these mediators increased with tumor progression. The effect of

treatment

with the B-E8 mAb on the tumor progression and survival was

evaluated.

This treatment significantly slowed down the tumor development:

on day 54,

there were more mice with low levels of sCD138 and hIL-10 in the

treated

group than in controls (p = 0.03 and 0.02, respectively). treatment

also

delayed death (median date of death was day 65 for control mice

and day 84

for anti-IL-6 mAb-treated mice; p < 0.02). Thus, hIL-6 is

expressed in

addition to vIL-6 in \*\*\*HHV\*\*\*-8-positive malignant

B

lymphocytes, and the viral cytokine does not totally substitute for

human

IL-6 in promoting tumor progression.

L16 ANSWER 2 OF 10 MEDLINE

AN 1999276344 MEDLINE

DN 99276344

TI The role of \*\*\*HHV\*\*\*-8 in Kaposi's sarcoma.

AU \*\*\*Neipel P\*\*\*; Fleckenstein B

CS Institut für Klinische und Molekulare Virologie, Universität

Erlangen-Nürnberg, Erlangen, D-91054, Germany.

SO SEMINARS IN CANCER BIOLOGY. (1999 Jun) 9 (3) 151-64.

Ref: 82

Journal code: A6Y ISSN: 1044-579X

CY United States

DT Journal Article; (JOURNAL ARTICLE)

General Review; (REVIEW)

(REVIEW, TUTORIAL)

LA English

FS Priority Journals

EM 199909

EW 19990901

AB The epidemiology of Kaposi's sarcoma (KS) amongst North

American and

Northern European patients with AIDS suggests that an infectious

agent

other than HIV is involved in its pathogenesis. Several lines of

evidence

indicate that human herpesvirus 8 (\*\*\*HHV\*\*\*-8) also termed

Kaposi's sarcoma associated herpesvirus, is the sought after agent.

DNA of

\*\*\*HHV\*\*\*-8 is invariably found in all forms of KS

where the

virus is present in the KS spindle cell. In contrast, \*\*\*HHV\*\*\*-8

\*\*\*g\*\*\* DNA is not regularly detected in most other

malignancies.

Antibodies against \*\*\*HHV\*\*\*-8 are more frequently

found in

groups at risk of KS, and \*\*\*HHV\*\*\*-8 precedes KS development. Several

genes have been

identified that exhibit transforming potential in cell culture systems.

In

addition, the virus encodes and induces several cytokines and

angiogenic

factors. This is of particular interest as models of KS pathogenesis

developed before the discovery of \*\*\*HHV\*\*\*-8 emphasized the

importance of inflammatory cytokines. Although the expression

pattern of

viral genes in KS is not certain yet, it appears likely that the

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pathogenic role of \*\*\*HHV\*\*\*. \*\*\*\*g\*\*\*\* in KS may be rather complex and differs from other virus-induced malignancies. 1999 Academic Press.

L16 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2000 ACS  
AN 1998:89360 CAPLUS  
DN 128:166368

TI The \*\*\*interleukin\*\*\* 6 of human herpesvirus 8 and its use in diagnostics and therapeutics  
IN \*\*\*Fleckenstein, Bernhard\*\*\* ; \*\*\*Albrecht, Jens-Christian\*\*\* ;  
\*\*\*Neipel, Frank\*\*\* ; \*\*\*Friedman-Kien, Alvin\*\*\* ;  
\*\*\*Huang\*\*\*

\*\*\* Yao-Qi\*\*\*  
PA Behring Diagnostics G.m.b.H. Germany; New York University; Fleckenstein, Bernhard, Albrecht, Jens-Christian, Neipel, Frank, Friedman-Kien, Alvin.  
Huang, Yao-Qi  
SO PCT Int. Appl., 19 pp.  
CODEN: PDXD2

DT Patent  
LA English  
FANGNT 1  
PATENT NO. KIND DATE APPLICATION NO.  
DATE

P1 WO 9803657 A1 19980129 WO 1996-EP3199  
19960719

W. US

RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE  
EP 912742 A1 19990506 EP 1996-927558 19960719  
R: AT, BE, CH, DE, ES, FR, GB, IT, LI, LU, NL, SE  
PRAI WO 1996-EP3199 19960719

AB Human herpesvirus 8 is found to carry a gene for an \*\*\*interleukin\*\*\*  
6 that can bind to the \*\*\*interleukin\*\*\* 6 receptor. The \*\*\*interleukin\*\*\* and the gene encoding can be used in the diagnosis and

treatment of a no. of diseases including: Kaposi sarcoma, Castleman's disease, multiple myeloma, kidney cell carcinoma, mesangial proliferative glomerulonephritis or B cell lymphoma. The protein may be manifold by expression of the cloned gene.

L16 ANSWER 4 OF 10 MEDLINE DUPLICATE

AN 1998158619 MEDLINE

DN 98158619

TI \*\*\*Human\*\*\* \*\*\*herpesvirus\*\*\* \*\*\*type\*\*\*  
\*\*\*\*g\*\*\*\*

\*\*\*interleukin\*\*\* -6 homologue is functionally active on human myeloma

cells [see comments]

CM Comment in: Blood 1998 Sep 15;92(6):2186-8

AU Burger R, \*\*\*Neipel F\*\*\*, Fleckenstein B, Savino R, Ciliberto G, Kalden J R, Gramatzki M

CS Division of Hematology/Oncology, Department of Medicine III, the Institute for Clinical and Molecular Virology, University of Erlangen-Nuremberg.

Erlangen, Germany.

SO BLOOD (1998 Mar 15) 91 (6) 1858-63.

Journal code: A8G. ISSN: 0006-4971.

CY United States  
DT Journal, Article, (JOURNAL ARTICLE)

LA English

FS Abridged Index Medicus Journals, Priority Journals, Cancer Journals

EM 199806

AB Seropidemiology and polymerase chain reaction studies have strongly suggested that \*\*\*human\*\*\* \*\*\*herpesvirus\*\*\*

\*\*\*type\*\*\*

\*\*\*\*g\*\*\*\* ( \*\*\*HHV\*\*\*. \*\*\*\*g\*\*\*\* ) is associated with Kaposi's sarcoma,

Castleman's disease, and body cavity-based lymphoma. The genome of

\*\*\*HHV\*\*\*. \*\*\*\*g\*\*\*\* harbors a viral analogue of the \*\*\*interleukin\*\*\* -6 (IL-6) gene. The amino acid sequence of

the viral IL-6 (vIL-6) protein is 74.7% identical to human IL-6 (hIL-6). IL-6 as a

B-cell growth and differentiation factor is known to play an essential role in the pathophysiology of B-cell tumors. Thus, it seems

possible that virus-encoded IL-6 contributes to malignant growth of \*\*\*HHV\*\*\*.

\*\*\*\*g\*\*\*\* -positive B-cell lymphatic tumors. We have tested a preparation

of \*\*\*HHV\*\*\*. \*\*\*\*g\*\*\*\* -derived IL-6 for the ability to promote the

proliferation of the human myeloma cell line INA-6, which is strictly

dependent on exogenous IL-6 for growth and survival. Viral IL-6 significantly induced DNA synthesis of INA-6 cells, but required

much more protein on a weight basis when compared with hIL-6 for maximal proliferation. The proliferative effect of vIL-6 was almost

completely inhibited by a combination of anti-IL-6 receptor (IL-6R) and anti-gp130

antibodies or IL-6R superantagonist Smt7 and anti-gp130 antibodies. This

report demonstrates that vIL-6 has proliferative activity on human cells

and that the IL-6R and gp130 are involved in vIL-6 signaling in the myeloma cell line INA-6.

L16 ANSWER 5 OF 10 BIOSIS COPYRIGHT 2000 BIOSIS

AN 1999:8773 BIOSIS

DN PREV199900087733

TI \*\*\*Human\*\*\* \*\*\*herpesvirus\*\*\* \*\*\*type\*\*\*  
\*\*\*\*g\*\*\*\* (

\*\*\*HHV\*\*\*. \*\*\*\*g\*\*\*\* ) IL-6 homologue induces proliferation of a human

myeloma cell line: Involvement of the IL-6 receptor complex.

AU Burger, R.; \*\*\*Neipel, F.\*\*\* ; Savino, R.; Gramatzki, M.

CS Div. Hemato/Oncol., Dep. Med. III, Inst. Clin. Mol. Virol, Univ.

Erlangen-Nuremberg, Erlangen Germany

SO Annals of Hematology (1998) Vol. 77, No. SUPPL. 2, pp. S103.

Meeting Info.: Annual Congress of the German and Austrian Societies of

Hematology and Oncology Frankfurt, Germany October 25-28, 1998 Austrian

Society of Hematology and Oncology

. ISSN: 0939-5555.

DT Conference

LA English

L16 ANSWER 6 OF 10 MEDLINE DUPLICATE

AN 1998374969 MEDLINE

DN 98374969

TI Human herpesvirus 8--the first human Rhabdovirus.

AU \*\*\*Neipel F\*\*\* ; Albrecht J C; Fleckenstein B

CS Institut für Klinische und Molekulare Virologie, Universitat Erlangen-Nürnberg, Germany.

Fleckenstein@viro.med.uni-erlangen.de

SO JOURNAL OF THE NATIONAL CANCER INSTITUTE.

MONOGRAPHS. (1998) (23) 73-7.

Ref: 22

Journal code: ATR. ISSN: 1052-6773.

CY United States

DT Journal, Article, (JOURNAL ARTICLE)

General Review, (REVIEW)

(REVIEW, TUTORIAL)

LA English

FS Priority Journals

EM 199811

EW 19981102

AB Kaposi's sarcoma (KS)-associated herpesvirus, also known as human herpesvirus 8 ( \*\*\*HHV\*\*\*. \*\*\*\*g\*\*\*\* ), is the first known human member

of the genus Rhabdovirus. It is regularly found by polymerase chain reaction in all forms of KS, in certain types of Castleman's disease, and in body cavity-based B-cell lymphoma. Other members of this virus group occur in nonhuman primates, ungulates, rabbits, and mice and cause fulminant lymphomas and other neoplastic disorders of the

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hematopoietic system. Rhadinoviruses share a typical genome structure, most characteristically, they contain numerous sequences that appear to be sequestered from cellular DNA. We cloned and sequenced almost the complete genome of \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* from a single K5 biopsy specimen. Although this procedure revealed collinear organization and extensive homologies with the open reading frames of herpesvirus saimiri, genes with homology to the known oncoproteins (Sip, Tip) were not identified in the \*\*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* genome. However, \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* reading frame K1, the positional analogue of Sip/Tip, was found to be significantly variable between different strains. We found, in addition, the reading frames for homologues of cellular \*\*\*interleukin\*\*\* macrophage inflammatory proteins alpha and beta (MIP1 alpha and MIP1 beta, respectively), an interferon-responsive factor, and two inhibitors of apoptosis. Several of these cell-homologous genes of \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* have already been shown to code for functional proteins.

L16 ANSWER 7 OF 10 MEDLINE DUPLICATE  
 4 AN 97138401 MEDLINE  
 DN 97138401  
 TI Human herpesvirus 8 encodes a homolog of \*\*\*interleukin\*\*\*  
 6. \*\*\*Neipel F\*\*\* ; Albrecht J C; Ensser A; Huang Y Q, Li J J, \*\*\*Friedman-Kien A E\*\*\* ; Fleckenstein B  
 CS Institut für Klinische und Molekulare Virologie, Universität Erlangen-Nürnberg, Erlangen, Germany.  
 neipel@viro.med.uni-erlangen.de  
 SO JOURNAL OF VIROLOGY, (1997 Jan) 71 (1) 839-42.  
 Journal code: KCV. ISSN: 0022-538X.  
 CY United States  
 DT Journal Article, (JOURNAL ARTICLE)  
 LA English  
 FS Priority Journals; Cancer Journals  
 OS GENBANK-U93872  
 EM 199704  
 AB Kaposi's sarcoma is a multifocal lesion that is reported to be greatly influenced by cytokines such as \*\*\*interleukin\*\*\* -6 (IL-6) and oncostatin M. DNA sequences of a novel human gammaherpesvirus, termed human herpesvirus 8 ( \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* ) or Kaposi sarcoma-associated herpesvirus, have been identified in all epidemiological forms of Kaposi's sarcoma with high frequency. The

presence of \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* DNA is also clearly associated with certain B-cell lymphomas (body cavity-based lymphomas) and multicentric Castleman's disease. Sequence analysis of a 1.7 kb fragment revealed that adjacent to a block of conserved herpesvirus genes (major DNA-binding protein, glycoprotein B, and DNA polymerase), the genome of \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* encodes structural homolog of IL-6. This cytokine is involved not only in the pathogenesis of Kaposi's sarcoma but also in certain B-cell lymphomas and multicentric Castleman's disease. The viral counterpart of IL-6 (vIL-6) has conserved important features such as cysteine residues involved in disulfide bridging or an amino-terminal signal peptide. Most notably, the region known to be involved in receptor binding is highly conserved in vIL-6. This conservation of essential features and the remarkable overlap between diseases associated with \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* and diseases associated with IL-6 clearly suggest that vIL-6 is involved in \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* pathogenesis.

L16 ANSWER 8 OF 10 BIOSIS COPYRIGHT 2000 BIOSIS  
 AN 1997:379203 BIOSIS  
 DN PREV19979678406  
 TI \*\*\*Human\*\*\* \*\*\*herpesvirus\*\*\* \*\*\*type\*\*\*  
 \*\*\*\*g\*\*\*\* ( \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* ) \*\*\*interleukin\*\*\* -6 homolog is functionally active on human plasma cells.  
 AU Burger, R. (1); \*\*\*Neipel, F.\*\*\* ; Fleckenstein, B.; Gramatzki, M.  
 CS (1) Div. Hematology/Oncol., Dep. Med. III, Univ. Erlangen-Nürnberg, D-91054 Erlangen Germany  
 SO Journal of Molecular Medicine (Berlin), (1997) Vol. 75, No. 7, pp. B184.  
 Meeting Info.: XIX Symposium of the International Association for Comparative Research on Leukemia and Related Diseases Heidelberg, Germany July 13-18, 1997  
 ISSN: 0946-2716.  
 DT Conference, Abstract  
 LA English

L16 ANSWER 9 OF 10 BIOSIS COPYRIGHT 2000 BIOSIS  
 AN 1998:66687 BIOSIS  
 DN PREV19980066687  
 TI \*\*\*Human\*\*\* \*\*\*herpesvirus\*\*\* \*\*\*type\*\*\*  
 \*\*\*\*g\*\*\*\* (

\*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* ) \*\*\*interleukin\*\*\* -6 homologue is functionally active on human myeloma cells.  
 AU Gramatzki, M.; Burger, R.; \*\*\*Neipel, F.\*\*\* ; Fleckenstein, B.  
 CS Div. Hematology/Oncology, Dep. Med. III, Univ. Erlangen-Nürnberg, D-91054 Erlangen Germany  
 SO Blood, (Nov. 15, 1997) Vol. 90, No. 10 SUPPL. 1 PART 1, pp. 87A.  
 Meeting Info.: 39th Annual Meeting of the American Society of Hematology San Diego, California, USA December 5-9, 1997 The American Society of Hematology  
 ISSN: 0006-4971.  
 DT Conference  
 LA English

L16 ANSWER 10 OF 10 BIOSIS COPYRIGHT 2000 BIOSIS  
 AN 1997:327181 BIOSIS  
 DN PREV199799626384  
 TI Does \*\*\*HHV\*\*\*\*. \*\*\*\*g\*\*\*\* play a role in the pathogenesis of Kaposi's sarcoma (KS)? The study of the expression of vIL-6 and v-cyclin D in KS tumor lesions.  
 AU \*\*\*Huang, Yao-qi (1)\*\*\* ; Li, Jian-Jun (1); \*\*\*Neipel, Frank\*\*\* ; Nicolaides, Alexander; \*\*\*Fleckenstein, Bernhard\*\*\* ; \*\*\*Friedman-Kien, Alvin E. (1)\*\*\*  
 CS (1) Dep. Microbiol., NYU Med. Center, Erlangen Germany  
 SO Journal of Acquired Immune Deficiency Syndromes and Human Retrovirology, (1997) Vol. 14, No. 4, pp. A25.  
 Meeting Info.: National AIDS Malignancy Conference Bethesda, Maryland, USA April 28-30, 1997  
 ISSN: 1077-9430.  
 DT Conference, Abstract  
 LA English

=> s (herpesvirus type 8 or human herpesvirus type 8 or herpes virus type 8 or hhv-8)/ab,bi  
 'AB IS NOT A VALID FIELD CODE  
 L17 1800 (HERPESVIRUS TYPE 8 OR HUMAN  
 HERPESVIRUS TYPE 8 OR HERPES VIRUS  
 TYPE 8 OR HHV-8)/AB,BI  
 => s (IL7 and interleukin-6)/ab,bi  
 'AB IS NOT A VALID FIELD CODE  
 L18 124 IL7 AND INTERLEUKIN-6/AB,BI  
 => s (IL7 and (viral interleukin-6 or v-il-6)/ab,bi

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AB IS NOT A VALID FIELD CODE  
L19 29 L17 AND (VIRAL INTERLEUKIN-6 OR  
VIL-6)/AB,BI

=> dup rem 119

PROCESSING COMPLETED FOR L19  
L20 11 DUP REM L19 (18 DUPLICATES REMOVED)

=> d 1- bib ab

YOU HAVE REQUESTED DATA FROM 11 ANSWERS -  
CONTINUE? Y(N)Y

L20 ANSWER 1 OF 11 BIOSIS COPYRIGHT 2000 BIOSIS  
AN 2000.324907 BIOSIS  
DN PREV20000324907

T1 Human herpesvirus 8-encoded interleukin-6 homologue (viral  
IL-6) induces  
endogenous human IL-6 secretion.  
AU Mori, Yasuko (1); Nishimoto, Norihito; Ohno, Mitsu; Inagi,  
Reiko;  
Diepckson, Penadda; Annou, Kiyoko; Yoshizaki, Kazuyuki;  
Yamanishi, Koichi  
CS (1) Department of Microbiology, Osaka University Medical  
School, Osaka  
University, 2-2 Yamadaoka, Suita, Osaka, 565-0871 Japan  
SO Journal of Medical Virology, (July, 2000) Vol. 61, No. 3, pp.  
332-335.  
print.  
ISSN: 0146-6615.

DT Article  
LA English  
SL English

AB We found that human herpesvirus 8-encoded IL-6 (VIL-6)  
induced endogenous  
human IL-6 (hIL-6) secretion from various cell lines (M1-4,  
THP-1, U937,  
Raji, and CESS) including patients with multicentric Castlemans  
disease.  
Especially, in M1-4 cells, hIL-6 was enhanced with VIL-6 by  
30-fold

compared with that of control. In addition, reverse transcriptase-  
polymerase chain reaction confirmed that VIL-6 induced hIL-6  
expression  
in M1-4 cells. Our novel finding of the hIL-6 induction by VIL-6  
indicates that VIL-6 may play a significant role in the pathogenesis  
of

\*\*\*HHV\*\*\* - \*\*\*g\*\*\* associated diseases.

L20 ANSWER 2 OF 11 MEDLINE DUPLICATE

AN 1999214337 MEDLINE  
DN 99214337

T1 Cellular tropism and \*\*\*viral\*\*\* \*\*\*interleukin\*\*\* -  
\*\*\*g\*\*\*

expression distinguish human herpesvirus 8 involvement in  
Kaposi's  
sarcoma, primary effusion lymphoma, and multicentric Castlemans  
disease.

AU Staskus K A; Sun R; Miller G; Racz P; Jaskowski A; Metricka C;  
Brett-Smith  
H; Haase A T  
CS Department of Microbiology, University of Minnesota Medical  
School,  
Minneapolis, Minnesota 55455, USA.  
kathryn@lenti.med.umn.edu  
NC CA-75172 (NCI)  
CA-70036 (NCI)

SO JOURNAL OF VIROLOGY, (1999 May) 73 (5) 4181-7.  
Journal code: KCV. ISSN: 0022-538X.  
CY United States

DT Journal, Article, (JOURNAL ARTICLE)  
LA English

FS Priority Journals; Cancer Journals  
EM 199907  
EW 19990704  
AB Human herpesvirus 8 ( \*\*\*HHV\*\*\* - \*\*\*g\*\*\* ) infection has  
been  
implicated in the etiology of Kaposi's sarcoma (KS), primary  
effusion  
lymphoma (PEL), and multicentric Castlemans disease (MCD),  
three diseases  
that frequently develop in immunocompromised, human  
immunodeficiency  
virus-positive individuals. One hypothesis that would account for  
different pathological manifestations of infection by the same virus  
is

that viral genes are differentially expressed in heterogeneous cell  
types.  
To test this hypothesis, we analyzed the localization and levels of  
expression of two viral genes expressed in latent and lytic  
infections and  
the viral homologue of interleukin-6 (VIL-6). We show that PEL  
parallels  
KS in the pattern of latent and lytic cycle viral gene expression but  
that  
the predominant infected cell type is a B cell. We also show that  
MCD  
differs from KS not only in the infected cell type (B-cell and T-cell  
lineage) but also in the pattern of viral gene expression. Only a few  
cells in the lesion are infected and all of these cells express  
lytic-cycle genes. Of possibly greater significance is the fact that in  
a  
comparison of KS, PEL, and MCD, we found dramatic differences  
in the  
levels of expression of VIL-6. Interleukin-6 is a B-cell growth and  
differentiation factor whose altered expression has been linked to  
plasma  
cell abnormalities, as well as myeloid and lymphoid malignancies.  
Our  
findings support the hypothesis that \*\*\*HHV\*\*\* - \*\*\*g\*\*\*  
plays an

important role in the pathogenesis of PEL and MCD, in which  
VIL-6 acts as  
an autocrine or paracrine factor in the lymphoproliferative processes  
common to both.

L20 ANSWER 3 OF 11 MEDLINE DUPLICATE

AN 1999290757 MEDLINE  
DN 99290757

T1 Angiogenesis and hematopoiesis induced by Kaposi's  
sarcoma-associated  
herpesvirus-encoded interleukin-6 [see comments].  
CM Comment in: Blood 1999 Jun 15;93(12):4031-3  
AU Aoki Y; Jaffe E S; Chang Y; Jones K; Tenen-Feldstein J; Moore  
P S; Tostato  
G

CS Division of Hematologic Products, Center for Biologics  
Evaluation and  
Research, Food and Drug Administration, Bethesda, MD, USA.

AOKI@CBER.FDA.GOV  
SO BLOOD, (1999 Jun 15) 93 (12) 4034-43.  
Journal code: ABG. ISSN: 0006-4971.

CY United States  
DT Journal, Article, (JOURNAL ARTICLE)

LA English  
FS Abridged Index Medicus Journals; Priority Journals; Cancer  
Journals

EM 199909  
EW 19990901

AB Kaposi's sarcoma-associated herpesvirus (KSHV, also known as  
human  
herpesvirus 8 [ \*\*\*HHV\*\*\* - \*\*\*g\*\*\* ]) is a herpesvirus  
linked to the  
development of Kaposi's sarcoma (KS), primary effusion  
lymphoma, and a  
proportion of Castlemans disease. KSHV encodes \*\*\*viral\*\*\*  
\*\*\*interleukin\*\*\* - \*\*\*g\*\*\* (VIL-6), which is structurally  
homologous  
to human and murine IL-6. The biological activities of VIL-6 are  
largely  
unknown. To gain insight into the biology of VIL-6, we expressed  
VIL-6 in  
murine fibroblasts NIH3T3 cells and inoculated stable  
VIL-6-producing  
clones into athymic mice. VIL-6 was detected selectively in the  
blood of  
mice injected with VIL-6-expressing clones. Compared with  
controls,  
VIL-6-positive mice displayed increased hematopoiesis in the  
myeloid,  
erythroid, and megakaryocytic lineages, plasmacytosis in spleen and  
lymph  
nodes, hepatosplenomegaly, and polyclonal  
hypergammaglobulinemia.

VIL-6-expressing NIH3T3 cells gave rise to tumors more rapidly  
than did  
control cells, and VIL-6-positive tumors were more vascularized

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than controls. Vascular endothelial growth factor (VEGF) was detected at higher levels in the culture supernatant of VIL-6-expressing cells compared with controls, and immunohistochemical staining detected VEGF in spleen, lymph nodes, and tumor tissues from mice bearing VIL-6-producing tumors but not control tumors. Thus, VIL-6 is a multifunctional cytokine that promotes hematopoiesis, plasmacytosis, and angiogenesis. Through these functions, VIL-6 may play an important role in the pathogenesis of certain KSHV-associated disorders.

L20 ANSWER 4 OF 11 MEDLINE DUPLICATE

AN 1999441800 MEDLINE  
DN 99441800

TI Human herpesvirus 8-encoded interleukin 6 activates HIV-1 in the UI monocytic cell line.

AU Gage J R, Breen E C, Echeverri A, Magganti L, Kishimoto T, Miles S, Martinez-Maza O

CS Department of Obstetrics and Gynecology, Osaka University, Japan.

NC CA57152 (NCI)  
CA73475 (NCI)  
F32CA80610 (NCI)

SO AIDS. (1999 Oct 1) 13 (14) 1851-5.  
Journal code: AID. ISSN: 0269-9370.

CY ENGLAND: United Kingdom

DI Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 200002

EW 20000204

AB OBJECTIVE: Human herpesvirus 8 ( \*\*\*HHV\*\*\* . \*\*\*g\*\*\* ) encodes a \*\*\*viral\*\*\* \*\*interleukin\*\*\* \*\*g\*\*\* (VIL-6) which is structurally and functionally similar to human interleukin 6 (hIL-6). Since hIL-6 has been shown to upregulate the expression of HIV-1, the objectives of this study were to examine the ability of VIL-6 to upregulate HIV-1, and to determine the interactions of this cytokine (viral cytokine) with the components of the interleukin 6 (IL-6) receptor

complex. DESIGN AND METHODS: Recombinant \*\*\*HHV\*\*\* . \*\*\*g\*\*\* VIL-6 (VIL-6) was assayed for bioactivity in the IL-6-dependent cell line MHH6 BSP2. HIV-1 p24 production by the UI monocytic and ACH-2 T-cell lines, which are chronically infected with HIV-1, was used to assess the

ability of VIL-6 to affect HIV-1 expression. hIL-6 and VIL-6 receptor utilization was determined by quantifying HIV-1 p24 production after neutralization of components of the IL-6 receptor complex, CD126L-6R, and CD130gp130, on UI cells with blocking antibodies. RESULT: \*\*\*HHV\*\*\* . \*\*\*g\*\*\* rVIL-6 was seen to have IL-6-like bioactivity in MHH6 BSP2 cells, and readily upregulated HIV-1 p24 production in UI monocytic cells, but not in ACH-2 T cells. The VIL-6 appeared to utilize the IL-6-specific component of the IL-6 signaling complex, CD126L-6R.

CONCLUSIONS: \*\*\*HHV\*\*\* . \*\*\*g\*\*\* VIL-6 clearly has the potential to upregulate HIV-1 expression in monocytic cells, and therefore may play a role in AIDS pathogenesis in individuals infected with both viruses.

L20 ANSWER 5 OF 11 MEDLINE DUPLICATE

AN 1999367590 MEDLINE  
DN 99367590

TI Heterogeneity of viral IL-6 expression in \*\*\*HHV\*\*\* . \*\*\*g\*\*\* -associated diseases

AU Cannon J S, Nicholas J, Orenstein J M, Mann R B, Murray P G, Browning P J, DiGiuseppe J A, Ceszmann E, Hayward G S, Ambinder R F

CS Department of Pharmacology, Johns Hopkins University School of Medicine.

Baltimore, Maryland, USA

NC UOI CA70062 (NCI)  
PO30 CA06973 (NCI)  
PO30 CA68485 (NCI)

SO JOURNAL OF INFECTIOUS DISEASES. (1999 Sep) 180 (3) 824-8.  
Journal code: JH3. ISSN: 0022-1899.

CY United States

DI Journal; Article; (JOURNAL ARTICLE)

LA English

FS Abstracted Index Medicus Journals; Priority Journals

EM 199911

EW 19991105

AB In order to characterize the expression of the \*\*\*viral\*\*\* \*\*interleukin\*\*\* \*\*g\*\*\* (VIL-6) homologue in various human herpesvirus 8 ( \*\*\*HHV\*\*\* . \*\*\*g\*\*\* )-associated diseases, in situ hybridization and immunohistochemistry were applied to formalin-fixed specimens. These assays showed consistent expression of VIL-6 in

primary effusion lymphomas and in a case of human immunodeficiency virus (HIV)-associated lymphadenopathy with a Castleman's disease-like appearance. In contrast, Kaposi's sarcoma specimens showed marked differences among specimens. In a consecutive series of specimens from the Johns Hopkins archives, VIL-6 expression was demonstrated in one of 13 cases. However, among 7 specimens selected from the AIDS Malignancy Bank because of their high levels of the T1.1 lytic transcript and virion production, VIL-6 expression was consistently demonstrated in infiltrating mononuclear cells and occasional spindle-shaped cells. Thus VIL-6 expression in clinical specimens correlates with other measures of the lytic viral cycle. Both assays generally give congruent results and are consistent with the possibility that VIL-6 expression plays a role in the pathogenesis of a variety of \*\*\*HHV\*\*\* . \*\*\*g\*\*\* -associated diseases.

L20 ANSWER 6 OF 11 MEDLINE

AN 1999399397 MEDLINE  
DN 99399397

TI Expression of human herpesvirus 8 ( \*\*\*HHV\*\*\* . \*\*\*g\*\*\* ) encoded pathogenic genes in Kaposi's sarcoma (KS) primary lesions.

AU Aschert G, Hohenadl C, Monini P, Zietz C, Browning P J, Ensoli B, Sturz M

CS Max Planck Institute for Biochemistry, Department of Virology, Martinsried, Germany.

SO ADVANCES IN ENZYME REGULATION. (1999) 39 331-9.  
Journal code: ZLG. ISSN: 0065-2571.

CY ENGLAND: United Kingdom

DI Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 200001

EW 20000104

AB Transcription of six different \*\*\*HHV\*\*\* . \*\*\*g\*\*\* specific mRNAs was examined in early- and late-stage KS primary lesions. Expression of the latency-associated T0.7 mRNA and of VP23 mRNA which is a specific marker of lytic/productive infection suggested that \*\*\*HHV\*\*\* . \*\*\*g\*\*\* is secondarily recruited into the KS lesions by productively infected monocytes/macrophages. From these cells \*\*\*HHV\*\*\* . \*\*\*g\*\*\* is transcribed to the KS spindle cells, which are latently infected. v-BCL-2, v-MCP-1 and \*\*\*v\*\*\* . \*\*\*IL\*\*\* . \*\*\*g\*\*\* were not

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expressed in latently infected KS spindle cells, therefore the impact of these factors in KS pathogenesis appears to be low. By contrast, v-Cyclin D was highly expressed in almost all latently infected spindle cells and may therefore be an important factor triggering progression of late-stage KS lesions.

L20 ANSWER 7 OF 11 MEDLINE DUPLICATE

AN 199913187 MEDLINE

TI Induction of human herpesvirus-8 DNA replication and transcription by butyrate and TPA in BCBL-1 cells.

AU Yu Y; Black J B; Goldsmith C S; Browning P J; Bhalla K; Offermann M K

CS Winship Cancer Center, Emory University, Atlanta, GA 30322, USA.

NC R01 CA67382 (NCI) P30AR42687 (NIAMS)

SO JOURNAL OF GENERAL VIROLOGY, (1999 Jan) 80 (Pt 1) 83-90.

Journal code: 19B ISSN: 0022-1317.

CY ENGLAND: United Kingdom DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals; Cancer Journals EM 1999005

EW 19990052 AB Human herpesvirus-8 (\*\*\*HHV\*\*\* - \*\*\*g\*\*\*) is a gammaherpesvirus that

is present primarily in a state of low level persistence in primary effusion lymphoma cell lines. Using BCBL-1 cells that harbour \*\*\*HHV\*\*\*

- \*\*\*g\*\*\* but lack Epstein-Barr virus, we demonstrate that sodium

butyrate is much more effective than the phorbol ester 12-O-tetradecanoyl

phorbol-13-acetate (TPA) at inducing high levels of class II and III virus

transcription and viral DNA replication, but also initiates apoptosis. Apoptosis occurs prior to assembly of virions when high

concentrations of butyrate (1-3 mM) are used, whereas reduction of butyrate concentration to

0.3 mM decreases the rate of apoptosis and results in production and

secretion of enveloped virions that are visualized at high number by electron microscopy in approximately 20% of BCBL-1 cells

Butyrate induces much higher levels of multiple class II and class III transcripts than does TPA, including v-MIP1, \*\*\*v\*\*\* - \*\*\*II\*\*\* - \*\*\*6\*\*\* v-Bcl-2.

VGPCR and ORF26. A decrease in concentration of butyrate from 3

to 0.3 mM delays the peak induction of these genes, but peak levels remain higher than peak levels in response to TPA. These studies indicate that the massive apoptosis induced by 3 mM butyrate could be diminished and delayed by reduction of butyrate concentration to 0.3 mM, thereby allowing expression of high levels ofytic-associated genes and production of high yields of \*\*\*HHV\*\*\* - \*\*\*g\*\*\* virions.

L20 ANSWER 8 OF 11 BIOSIS COPYRIGHT 2000 BIOSIS

AN 1999379732 BIOSIS

DN PREVI99900379732 TI KSHV/ \*\*\*HHV\*\*\* - \*\*\*g\*\*\* interleukin-6 (vIL6) expression in

HIV-related lymphadenopathy correlates with development of Kaposi's

sarcoma and survival.

AU Chaddam, A. (1); Hyjek, E. (1); Ying, L. (1); Mulligan, L. (1); Knowles, D. M. (1); Cesman, E. (1)

CS (1) Weill Medical College-Cornell University, New York, NY USA

SO J AIDS Journal of Acquired Immune Deficiency Syndromes, (May 1, 1999) Vol.

21, No. 1, pp. A21.

Meeting Info.: Third National AIDS Malignancy Conference Bethesda, Maryland, USA May 26-27, 1999

DT Conference

LA English

L20 ANSWER 9 OF 11 CAPLUS COPYRIGHT 2000 ACS

AN 199889360 CAPLUS

DN 128166368 TI The interleukin 6 of human herpesvirus 8 and its use in

diagnostics and therapeutics

IN Fleckenstein, Bernhard; Albrecht, Jens-Christian; Neipel, Frank; Friedman-Kien, Alvin; Huang, Yao-Qi

PA Behring Diagnostics G.m.b.H., Germany; New York University; Fleckenstein, Bernhard; Albrecht, Jens-Christian; Neipel, Frank; Friedman-Kien, Alvin;

Huang, Yao-Qi

SO PCT Int Appl, 19 pp. CODEN: PIXXD2

DT Patent

LA English

FAN CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9803657 A1 19980129 WO 1996-EP3199

19960719 W: US

RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE

EP 912742 A1 19990306 EP 1996-977558 19960719

R: AT, BE, CH, DE, ES, FR, GB, IT, LI, LU, NL, SE

PRAI WO 1996-EP3199 19960719

AB Human herpesvirus 8 is found to carry a gene for an interleukin 6 that can

bind to the interleukin 6 receptor. The interleukin and the gene encoding

can be used in the diagnosis and treatment of a no. of diseases including: Kaposi sarcoma, Castlemans disease, multiple myeloma, kidney

cell carcinoma, mesangial proliferative glomerulonephritis or B cell lymphoma.

The protein may be manul. by expression of the cloned gene.

L20 ANSWER 10 OF 11 MEDLINE DUPLICATE

AN 97184526 MEDLINE

DN 97184526 TI A single 13-kilobase divergent locus in the Kaposi

sarcoma-associated herpesvirus (human herpesvirus 8) genome contains nine open

reading frames that are homologous to or related to cellular proteins.

AU Nicholas J; Ruvolo V; Zeng J; Cuido D; Guo H G; Reitz M S; Hayward G S

CS Department of Oncology, Johns Hopkins University School of Medicine, Baltimore, Maryland 21231, USA

NC U01 CA37314 (NCI) R01 CA73585 (NCI)

P30 CA06973 (NCI) SO JOURNAL OF VIROLOGY, (1997 Mar) 71 (3) 1963-74.

Journal code: KCV ISSN: 0022-538X

CY United States DT Journal; Article; (JOURNAL ARTICLE)

LA English FS Priority Journals; Cancer Journals

OS GENBANK-U67774; GENBANK-U67775; GENBANK-U74585; GENBANK-U83347; GENBANK-U83348; GENBANK-U83349; GENBANK-U83350; GENBANK-U83351; GENBANK-U83269

EM 199705 AB Two small fragments of a novel human gammaherpesvirus genome known as

Kaposi's sarcoma (KS)-associated herpesvirus or human herpesvirus 8 (\*\*\*HHV\*\*\* - \*\*\*g\*\*\*) have been shown to be present in

virtually all AIDS and non-AIDS KS lesions, as well as in body cavity-based lymphomas

(BCBL) and in multicentric Castlemans disease. We have

extended those studies by identifying and sequencing a third fragment of

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\*\*\*HHV\*\*\*  
 \*\*\*\*\* DNA encoding a viral thymidylate synthetase (TS) gene. Use of this viral TS fragment as a probe led to the identification and mapping of a cluster of overlapping phage lambda clones from a BCB1 tumor DNA genomic library that spanned 48 kb on the left-hand side of the \*\*\*HHV\*\*\*  
 \*\*\*\*\* genome between the equivalents of open reading frame 6 (ORF6) and ORF31 of herpesvirus saimiri (HVS). DNA sequencing of a 17-kb segment encompassing a gammaherpesvirus divergent locus (DL-B) between ORF11 and ORF17 revealed the presence of nine viral ORFs with predicted products related to cellular proteins. These include the complete TS gene and a dihydrofolate reductase (DHFR) gene, four novel cytokine genes (encoding \*\*\*viral\*\*\* \*\*interleukin\*\*\*. \*\*\*g\*\*\*, viral MIP-1A, viral MIP-1B, and BCK) that have not previously been found to be encoded by a virus, and a bcl-2 homolog. This region in \*\*\*HHV\*\*\*  
 \*\*\*\*\* also contains the T1.1 abundant lytic cycle nuclear RNA gene and encompasses two genes (or exons) encoding proteins with C4HC3 zinc finger domains of the PHD/leukemia-associated protein subtype. The latter are related to the spliced immediate-early IE1 protein of the gamma-2 class herpesvirus bovine herpesvirus type 4 and a similar motif found in HVS  
 ORF12. Although genes for TS and DHFR enzymes are also encoded by HVS (ORF70 and ORF2), both occur at different genomic loci than in \*\*\*HHV\*\*\*  
 \*\*\*\*\* and the \*\*\*HHV\*\*\*. \*\*\*g\*\*\* DHFR protein is much farther diverged from human DHFR than is the HVS version, implying that they were probably acquired as host cell cDNAs by independent evolutionary events. Transcripts from the IE1-A, IE1-B, DHFR, and MIP-1B genes were all detected by Northern blot hybridization analysis in a BCB1 cell line at 12 h after induction with butyrate but were not present before induction, indicating that these are all primarily lytic cycle genes. We conclude that the DL-B locus of gammaherpesviruses displays considerably more variability than previously appreciated and that expression of many

of these genes is likely to have important implications for \*\*\*HHV\*\*\*  
 \*\*\*\*\* biology and therapy.

L20 ANSWER 11 OF 11 BIOSIS COPYRIGHT 2000 BIOSIS AN 1997/327181 BIOSIS DN PRE V19799626384  
 TI Does \*\*\*HHV\*\*\*. \*\*\*g\*\*\* play a role in the pathogenesis of Kaposi's sarcoma (KS). The study of the expression of vIL-6 and v-glycin D in KS tumor lesions.  
 AU Huang Yao-Chi (1), Li Jian-Jun (1), Neipel, Frank; Nicolaides, Alexander, Fleckenstein, Bernhard; Friedman-Kien, Alvin E. (1)  
 CS (1) Dep. Microbiol., NYU Med. Center, Erlangen Germany  
 SO Journal of Acquired Immune Deficiency Syndromes and Human Retrovirology.  
 (1997) Vol. 14, No. 4, pp. A25  
 Meeting Info.: National AIDS Malignancy Conference Bethesda, Maryland, USA  
 April 28-30, 1997  
 ISSN: 1077-9450  
 DT Conference, Abstract  
 LA English

=> d his

(FILE HOME ENTERED AT 16:30:20 ON 15 SEP 2000)

FILE MEDLINE ENTERED AT 16:30:25 ON 15 SEP 2000  
 L1 14 S HERPES VIRUS TYPE 8/AB, BI  
 L2 1 S L1 AND INTERLEUKIN/AB, BI  
 L3 458 S HHV-8/AB, BI  
 L4 38 S L3 AND INTERLEUKIN/AB, BI  
 L5 16 S L4 AND IL-6/AB, BI  
 L6 57 S HUMAN HERPESVIRUS TYPE 8/AB, BI  
 L7 4 S L6 AND INTERLEUKIN/AB, BI

FILE MEDLINE, EMBASE, BIOSIS, INPADOC, CAPLUS  
 ENTERED AT 16:33:56 ON 15 SEP 2000

E FLECKENSTEIN BERNHARD/AU  
 L8 312 S E1-E10  
 E ALBRECHT JENS-CHRISTIAN/AU  
 E ALBRECHT JENS CHRISTIAN/AU  
 L9 51 S E1-E3  
 E NEIPEL FRANK/AU  
 L10 145 S E2-E4  
 E FRIEDMAN KIEN ALVIN/AU  
 L11 496 S E1-E4  
 E HUANG YAO QUAU  
 L12 48 S E1-E4  
 L13 937 S L8 OR L9 OR L10 OR L11 OR L12

L14 82 S L13 AND (L1 OR L3 OR L6)  
 L15 20 S L14 AND INTERLEUKIN/AB, BI  
 L16 10 DUP REM L15 (10 DUPLICATES REMOVED)  
 L17 1800 S (HERPESVIRUS TYPE 8 OR HUMAN  
 HERPESVIRUS TYPE 8 OR HERPES VIR  
 L18 124 S L17 AND INTERLEUKIN-6/AB, BI  
 L19 29 S L17 AND (VIRAL INTERLEUKIN-6 OR  
 V-IL-6)/AB, BI  
 L20 11 DUP REM L19 (18 DUPLICATES REMOVED)

=>

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 Executing the logoff script...

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FULL ESTIMATED COST ENTRY SESSION 153.92 159.81

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RB113.N37

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311794

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